

**METHODS FOR DECREASING NON-SPECIFIC BINDING
OF BEADS IN DUAL BEAD ASSAYS INCLUDING RELATED
OPTICAL BIODISCS AND DISC DRIVE SYSTEMS**

5

CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a continuation-in-part of U.S. Application Serial No. 09/997,741 filed November 27, 2001 which claimed the benefit of priority from U.S. Provisional Application Serial No. 60/253,283 filed November 27, 2000; U.S. Provisional Application Serial No. 60/253,958 filed November 28, 2000; and U.S. Provisional Application Serial No. 60/272,525 filed March 1, 2001.

This application also claims the benefit of priority from U.S. Provisional Application Serial No. 60/272,134 filed the February 28, 2001; U.S. Provisional Application Serial No. 60/272,243 also filed February 28, 2001; U.S. Provisional Application Serial No. 60/272,525 filed March 1, 2001; U.S. Provisional Application Serial No. 60/275,006 filed March 12, 2001; U.S. Provisional Application Serial No. 60/275,643 filed March 14, 2001; U.S. Provisional Application Serial No. 60/278,691 filed March 26, 2001; U.S. Provisional Application Serial No. 60/314,906 filed August 24, 2001; and U.S. Provisional Application Serial No. 60/352,270 filed January 30, 2002. Each of the above utility and provisional applications is herein incorporated by reference in its entirety.

BACKGROUND OF THE INVENTION**1. Field of the Invention**

The present invention relates to optical analysis discs and optical bio-discs. The invention further relates to methods for decreasing non-specific bindings of beads in bead assays and in particular in dual bead assays. This invention is also directed to dual bead assays performed on optical bio-discs and related drive systems and methods.

2. Discussion of the Related Art

There is a significant need to make diagnostic assays and forensic assays of all types faster and more local to the end-user. Ideally, clinicians, patients, investigators, the military, other health care personnel, and consumers should be able to test themselves for the presence of certain factors or indicators in their systems, and for the presence of certain biological material at a crime scene or on a

1

I certify that this document was submitted to the U.S. Patent and Trademark Office using U.S. Express Mail # EL923484765US
 Jacquelyn Campbell
 Print Name
Jacquelyn Campbell February 28, 2002
 Signature Date

10087549-022802

battlefield. At present, there are a number of silicon-based chips with nucleic acids and/or proteins attached thereto, which are commercially available or under development. These chips are not for use by the end-user, or for use by persons or entities lacking very specialized expertise and expensive equipment.

5

SUMMARY OF THE INVENTION

The present invention relates to performing assays, and particularly to using dual bead structures on a disc. The invention includes methods for preparing assays, methods for performing assays, discs for performing assays, and related
10 detection systems.

According to one aspect of the present invention, there is provided a method for identifying whether a target agent is present in a biological sample. The method includes the steps of preparing a plurality of capture beads pre-treated with a bead blocking agent, each of the capture beads having at least one transport probe
15 affixed thereto, preparing a plurality of reporter beads pre-treated with the bead blocking agent, each of the reporter beads having at least one signal probe affixed thereto, mixing the capture beads and the reporter beads under binding conditions so as to permit formation of a dual bead complex if the target agent is present in the sample, the reporter bead and capture bead each being bound to the target agent,
20 isolating the dual bead complex from the mixture to obtain an isolate, exposing the isolate to a capture field on a disc, the capture field having a capture agent that binds to the dual bead complex, and detecting the presence of the dual bead complex in the disc to indicate that the target agent is present in the sample. In one embodiment of this method, the bead blocking agent is a detergent. In another
25 embodiment thereof, the bead blocking agent is an anionic, cationic, or non-ionic molecule. And in a preferred embodiment, the bead blocking agent is sheared denatured salmon sperm DNA.

According to another aspect of this invention, the capture beads and the reporter beads are pretreated with the bead blocking agent for approximately one
30 hour at room temperature prior to target capture. In one preferred embodiment of this method, the capture beads are magnetic beads pre-incubated with the salmon sperm DNA to thereby block sites available for non-specific binding of the target agent.

These embodiments of method according to this aspect of the present invention may advantageously include the further step of treating the capture beads and reporter beads with a probe blocking agent prior to target capture. In one embodiment of this aspect of the present invention, the probe blocking agents employed are DNA sequences complementary to the capture probes so that the probe blocking agents bind to the transport probes to thereby reduce the density of available transport probes for target binding. In another embodiment thereof, the probe blocking agents employed are DNA sequences complementary to the reporter probes so that the probe blocking agents bind to the signal probes to thereby reduce the density of available signal probes for target binding.

In any of the above methods, the mixing step may be preferably performed in an intermittent manner.

According to yet another aspect of this invention, the method may include the further step of evaluating the non-specific binding of the dual bead assay in the presence of salt concentrations ranging from 0.1M up to 1M. In one particular preferred embodiment, the present method may advantageously include the additional step of using a wash buffer having 10mM EDTA.

In accordance with another principal aspect of this invention, there is provided a method of preparing a dual bead assay for use in an optical bio-disc. This particular method includes the steps of providing a mixture of capture beads that have transport probes covalently bound thereto, providing a mixture of reporter beads that have signal probes covalently bound thereto, blocking the mixture of capture beads with a bead blocking agent, blocking the mixture of reporter beads with the bead blocking agent, suspending the mixture of capture beads in a hybridization solution, adding to the mixture a target agent that hybridizes with the transport probes, adding to the mixture the reporter beads, allowing the signal probes to hybridize with the target agent to thereby form a dual bead complex including at least one capture bead and one reporter bead, separating the dual bead complex from unbound reporter beads, removing from the mixture the unbound reporter beads, and loading the mixture including the dual bead complex into an optical bio-disc for analysis.

In this method, the step of adding the target agent may be preferably performed before the step of adding the reporter beads. In one particular embodiment thereof, the target agent is a segment of genetic material. The

segment of genetic material may be a single strand of DNA, or alternatively include a portion of double stranded DNA. In yet another alternative embodiment, the segment of genetic material may be a single strand of RNA which may also include a portion of double stranded RNA.

- 5 According to one specific aspect of this particular method, the capture beads may be magnetic and then the separating step is performed by use of a magnet field. This magnetic field may be formed by a magnet or by an electromagnet. This method may also advantageously include the further steps of removing the hybridization solution for the mixture, washing the dual bead complex to purify the
10 mixture by further removing unbound material, and/or adding a buffer solution to the mixture.

- In accordance with yet another principal aspect of this invention, there is provided an alternate method of preparing a dual bead assay for use in an optical bio-disc. This alternate method includes the steps of providing a mixture of capture
15 beads having transport probes covalently attached thereto, providing a mixture of reporter beads that have signal probes covalently bound thereto, blocking the mixture of capture beads with a bead blocking agent, blocking the mixture of reporter beads with the bead blocking agent, suspending the mixture of capture beads in a hybridization solution, adding to the mixture a target agent that hybridizes
20 with the transport probes, allowing the transport probes to hybridize with the target agent to thereby form a hybridized partial complex including at least one capture bead, separating within the mixture the hybridized partial complex from unbound target agents, adding to the mixture reporter beads including signal probes covalently attached thereto, allowing the signal probes to hybridize with the target
25 agent to thereby form a dual bead complex including at least one capture bead and one reporter bead, separating the dual bead complex from unbound reporter beads, removing from the mixture the unbound reporter beads, and loading the mixture including the dual bead complex into an optical bio-disc for analysis.

- Similarly in this alternate method, the step of adding the target agent may be
30 preferably performed before the step of adding the reporter beads. In one particular embodiment thereof, the target agent is a segment of genetic material. The segment of genetic material may be a single strand of DNA, or alternatively include a portion of double stranded DNA. In yet another alternative embodiment, the

segment of genetic material may be a single strand of RNA that may also include a portion of double stranded RNA.

In accordance with one specific aspect of this particular alternate method, the capture beads may be magnetic and then the separating step is performed by use
 5 of a magnet field. As with the prior method, this magnetic field may be formed by a magnet or by an electromagnet. Similarly, this alternate method may also advantageously include the further steps of removing the hybridization solution for the mixture, washing the dual bead complex to purify the mixture by further removing unbound material, and/or adding a buffer solution to the mixture.

10 According to yet another principal aspect of the present invention, there is also provided a first method of testing for the presence of a target-DNA in a DNA sample by use of an optical bio-disc. This first method of testing DNA includes the steps of preparing a DNA sample to be tested for the presence of a target-DNA, preparing a plurality of reporter beads each having covalently attached thereto a
 15 plurality of strands of signal-DNA and an anchor agent (the target-DNA and the signal-DNA being complementary), preparing a plurality of capture beads each having covalently attached thereto a plurality of transport-DNA (the target-DNA and transport-DNA being complimentary), blocking the plurality of reporter beads and the plurality of capture beads with a bead blocking agent, mixing the DNA sample,
 20 the plurality of reporter beads, and the plurality of capture beads to thereby form a test sample (the transport-DNA and the signal-DNA being non-complimentary), allowing hybridization between the signal-DNA, any target-DNA, and transport-DNA existing in the DNA sample to thereby form a dual bead complex including at least one capture bead and one reporter bead, removing from the test sample reporter
 25 beads and capture beads that are not associated with the dual bead complex, depositing the test sample in a flow channel of an optical bio-disc which is in fluid communication with a target zone, the target zone including a plurality of capture agents each including an amino group that attaches to an active layer to immobilize the capture agents within the target zone, allowing any anchor agent to bind with the
 30 capture agents so that reporter beads associated with the dual bead complex are maintained within the target zone, and detecting any dual bead complexes in the target zone to thereby determine whether target-DNA is present in the DNA sample.

In accordance with still yet a further principal aspect of the present invention, there is provided a second method of testing for the presence of a target-DNA in a

test sample by use of an optical bio-disc. This second method of testing for a target-DNA includes the steps of preparing a test sample to be tested for the presence of a target-DNA, preparing a plurality of reporter beads each having covalently attached thereto a plurality of strands of signal-DNA, the target-DNA and the signal-DNA being complementary, preparing a plurality of capture beads each having covalently attached thereto a plurality of transport-DNA and an anchor agent, the target-DNA and transport-DNA being complimentary, blocking the plurality of reporter beads and the plurality of capture beads with a bead blocking agent, depositing a plurality of capture beads and reporter beads in a mixing chamber, each of the reporter beads and the capture beads including signal-DNA and transport-DNA, respectively, being non-complimentary to each other, depositing the test sample in the mixing chamber of an optical bio-disc which is linked to a target zone by a connecting flow channel allowing any target-DNA existing in the test sample to bind to the signal-DNA and the transport-DNA on the reporter and the capture bead, respectively, to thereby form a dual bead complex, rotating the optical bio-disc to cause the dual bead complex to move from the mixing chamber through the flow channel and into the target zone, the target zone including a plurality of capture agents each including an amino group that attaches to an active layer to immobilize the capture agents within the target zone, the capture agent having affinity for the anchor agent, allowing any anchor agent to bind with the capture agent so that capture beads associated with dual bead complex are maintained within the capture zone, removing from the target zone reporter beads that are free of any dual bead complex, and detecting any dual bead complex in the target zone to thereby determine whether target-DNA is present in the test sample.

The present invention also provides a method of testing for the presence of a target-RNA in a test sample by use of an optical bio-disc. This method of testing for target-RNA includes the steps of preparing a test sample to be tested for the presence of a target-RNA, preparing a plurality of reporter beads each having covalently attached thereto a plurality of strands of signal-DNA, the target-RNA and the signal-DNA being complementary, preparing a plurality of capture beads each having covalently attached thereto a plurality of transport-DNA and an anchor agent, the target-RNA and transport-DNA being complimentary, blocking the plurality of reporter beads and the plurality of capture beads with a bead blocking agent, depositing a plurality of capture beads and reporter beads in a mixing chamber,

each of the reporter beads and capture beads including the signal-DNA and the transport-DNA, respectively, being non-complimentary to each other, depositing the test sample in the mixing chamber of an optical bio-disc which is linked to a target zone by a connecting flow channel allowing any target-RNA existing in the test sample to hybridize with the signal-DNA and the transport-DNA on the reporter and the capture bead, respectively, to thereby form a dual bead complex, rotating the optical bio-disc to cause the dual bead complex to move from the mixing chamber through the flow channel and into the target zone, the target zone including a plurality of capture agents each including an amino group that attaches to an active layer to immobilize the capture agents within the target zone, the capture agent and the anchor agent having affinity to each other, allowing any anchor agent to bind with the capture agent so that capture beads associated with dual bead complex are maintained within the capture zone, removing from the target zone reporter beads that are free of any dual bead complex, and detecting any dual bead complex in the target zone to thereby determine whether target-RNA is present in the test sample.

According to yet another principal aspect of the present invention, there is provided a method of testing for the presence of a target-antigen in a test sample by use of an optical bio-disc. This method of testing for a target-antigen includes the steps of preparing a test sample to be tested for the presence of a target-antigen, preparing a plurality of reporter beads each having covalently attached thereto a plurality of signal-antibody, the signal-antibody having an affinity to epitopes on the target-antigen, preparing a plurality of capture beads each having covalently attached thereto a plurality of transport-antibody and an anchor agent, the transport-antibody having affinity to epitopes on the target-antigen, blocking the plurality of reporter beads and the plurality of capture beads with a bead blocking agent, depositing the capture beads and the reporter beads in a mixing chamber of an optical bio-disc, each of the reporter beads and capture beads including the signal-antibody and the transport-antibody, respectively, having no affinity to each other, depositing the test sample in the mixing chamber of the optical bio-disc which is linked to a target zone by a connecting flow channel allowing any target-antigen existing in the test sample to bind to the signal-antibody and the transport-antibody on the reporter and the capture bead, respectively, to thereby form a dual bead complex, rotating the optical bio-disc to cause the dual bead complex to move from the mixing chamber through the flow channel and into the target zone, the target

zone including a plurality of capture agents each including an amino group that attaches to an active layer to immobilize the capture agents within the target zone, allowing any anchor agent to bind with the capture agent so that capture beads associated with dual bead complex are maintained within the capture zone, removing from the target zone reporter beads that are free of any dual bead complex, and detecting any dual bead complex in the target zone to thereby determine whether target-antigen is present in the test sample.

In any of the above methods employing a bio-disc with a target zone, the dual bead complex may be detected by directing a beam of electromagnetic energy from a disc drive assembly toward the target zone and analyzing electromagnetic energy returned from the target zone. In addition in these methods, the bead blocking agent may preferably include sheared salmon sperm DNA.

Yet another aspect of this invention is to provide an optical bio-disc to be used in conjunction with any of the above methods. It is also an aspect of this invention to provide a disc drive assembly or system for operating or using

According to the manufacturing aspects of this invention, there is provided a method of making an optical bio-disc to test for the presence of a target agent in a test sample. This method of manufacturing includes the steps of providing a substrate having a center and an outer edge, encoding information on an information layer associated with the substrate, the encoded information being readable by a disc drive assembly to control rotation of the disc, forming a target zone in association with the substrate, the target zone disposed at a predetermined location relative to the center of the substrate, depositing an active layer in the target zone, depositing a plurality of capture agents in the target zone, each capture agent including an amino group that covalently attaches to the active layer to immobilize the capture agent within the target zone, and blocking the target zone with a plurality of blocking agents after depositing the capture agents. This disc manufacturing method also includes forming a flow channel in fluid communication with the target zone, forming a mixing chamber in fluid communication with the flow channel, and depositing a plurality of reporter beads in the mixing chamber. In this embodiment, each of the reporter beads has covalently attached thereto a plurality of signal probes, and each of the signal probes has an affinity to the target agent. The manufacturing method also includes the step of depositing a plurality of capture beads in the mixing chamber. Each of the capture beads has covalently attached

thereto, a plurality of transport probes and an anchor agent. Each of the transport probes has an affinity to the target agent. The transport probes and signal probes have no affinity toward each other. The capture agents and the anchor agents have specific affinity for each other. This disc manufacturing method also includes the

5 step of adding a pre-determined amount of bead blocking agent to the mixing chamber to prevent non-specific binding of the beads to each other and the walls of the mixing chamber.

More generally now, the present invention includes methods for determining whether a target agent is present in a biological sample. These methods can

10 include mixing capture beads, each having at least one transport probe, reporter beads, each having at least one signal probe, and a biological sample. These components are mixed under binding conditions that permit formation of a dual bead complex if the target agent is present in the sample. The dual bead complex thus includes a reporter bead and a capture bead each bound to the target agent. The

15 dual bead complex is isolated from the mixture to obtain an isolate. The isolate is then exposed to a capture field on an optical disc. The capture field has a capture agent that binds specifically to the signal probe or transport probe of the dual bead complex. The dual bead complex in the optical disc is then detected to indicate that the target agent is present in the sample and, if desired, to indicate a concentration.

20 The capture beads can have a specified size and have a characteristic that makes them "isolatable." The capture beads are preferably magnetic, in which case the isolating of dual bead complex (and some capture beads not part of a complex) in a mixture includes subjecting the mixture to a magnetic field with a permanent magnet or an electromagnet.

25 The reporter bead should have characteristics that make it identifiable and distinguishable with detection. The reporter beads can be made of one of a number of materials, such as latex, gold, plastic, steel, or titanium, and should have a known and specified size. The reporter beads can be fluorescent and can be yellow, green, red, or blue, for example.

30 The dual bead complex can be formed on the disc itself, or outside the disc and added to the disc. To form the dual bead complex off disc, methods referred to here as "single-step" or "two-step" can be employed. In the two-step method, the mixture initially includes capture beads and the sample. The capture beads are then isolated to wash away unbound sample and leave bound and unbound capture

beads in a first isolate. Reporter beads are then added to the first isolate to produce dual bead complex structures and the isolation process is repeated. The resulting isolate leaves dual bead complex with reporters, but also includes unbound capture beads without reporters. The reporters make the dual bead complex detectable.

- 5 In the "single-step" method, the capture beads, reporter beads, and sample are mixed together from the start and then the isolation process isolates dual bead complex along with unbound capture beads.

These methods for producing and isolating dual bead complex structures can be performed on the disc. The sample and beads can be added to the disc together, 10 or the beads can be pre-loaded on the disc so that only a sample needs to be added. The sample and beads can be added in a mixing chamber on the disc, and the disc can be rotated in one direction or in both to assist the mixing. An isolate can then be created, such as by applying an electromagnet and rotating to cause the material other than the capture beads to be moved to a waste chamber. The 15 isolate is then directed through rotation to capture fields.

The dual bead complex structures can be detected on the capture field by use of various methods. In one embodiment, the detecting includes directing a beam of electromagnetic energy from a disc drive toward the capture field and analyzing electromagnetic energy returned from or transmitted past the reporter 20 bead of the dual bead complex attached to the capture field. The disc drive assembly can include a detector and circuitry or software that senses the detector signal for a sufficient transition between light and dark (referred to as an "event") to spot a reporter bead.

Beads can, alternatively, be detected based on their fluorescence. In this 25 case, the energy source in the disc drive preferably has a wavelength controllable light source and a detector that is or can be made specific to a particular wavelength. Alternatively, a disc drive can be made with a specific light source and detector to produce a dedicated device, in which case the source may only need fine-tuning.

30 The biological sample can include blood, serum, plasma, cerebrospinal fluid, breast aspirate, synovial fluid, pleural fluid, peritoneal fluid, pericardial fluid, urine, saliva, amniotic fluid, semen, mucus, a hair, feces, a biological particulate suspension, a single-stranded or double-stranded nucleic acid molecule, a cell, an organ, a tissue, or a tissue extract, or any other sample that includes a target that

may be bound through chemical or biological processes. Further details relating to other aspects associated with the selection and detection of various targets is disclosed in, for example, commonly assigned and co-pending U.S. Provisional Patent Application Serial No. 60/278,697 entitled "Dual Bead Assays for Detecting Medical Targets" filed March 26, 2001, which is incorporated herein by reference in its entirety.

In addition to these medical uses, the embodiments of the present invention can be used in other ways, such as for testing for impurities in a sample, such as food or water, or for otherwise detecting the presence of a material, such as a biological warfare agent.

The target agent can include, for example, a nucleic acid (such as DNA or RNA) or a protein (such as an antigen or an antibody). If a nucleic acid, both the transport probe and the signal probe can be a nucleic acid molecule complementary to the target nucleic acid. If a protein, both the transport probe and the signal probe can be an antibody that specifically binds the target protein.

The transport probe or signal probe can bind specifically to the capture agent on the optical disc due to a high affinity between the probe and the capture agent. This high affinity can, for example, be the result of a strong protein-protein affinity (i.e., antigen-antibody affinity), or the result of a complementarity between two nucleic acid molecules.

Preferably the binding is to the signal probe, and then the disc is rotated to move unbound structures, including capture beads not bound to reporter beads, away from the capture field. If the binding is to the transport probe, unbound capture beads will be included, although the reporter beads are still the beads that are detected. This may be acceptable if the detection is for producing a yes/no answer, or if a fine concentration detection is not otherwise required.

The transport probe and signal probe can each be one or more probes selected from the group consisting of single-stranded DNA, double-stranded DNA, single-stranded RNA, peptide nucleic acid, biotin, streptavidin, an antigen, an antibody, a receptor protein, and a ligand. In a further embodiment, each transport probe includes double-stranded DNA and single-stranded DNA, wherein the double-stranded DNA is proximate to the capture layer of the optical disc and the single-stranded DNA is distal relative to the capture layer of the optical disc.

The reporter bead and/or signal probe can be biotinylated and the capture agent can include streptavidin or neutravidin. Chemistry for affixing capture agents to the capture layer of the optical disc are generally known, especially in the case of affixing a protein or nucleic acid to solid surfaces. The capture agent can be affixed
 5 to the capture layer by use of an amino group or a thiol group.

The target agent can include a nucleic acid characteristic of a disease, or a nucleotide sequence specific for a person, or a nucleotide sequence specific for an organism, which may be a bacterium, a virus, a mycoplasma, a fungus, a plant, or an animal. The target agent can include a nucleic acid molecule associated with
 10 cancer in a human. The target nucleic acid molecule can include a nucleic acid, which is at least a portion of a gene selected from the group consisting of *HER2neu*, *p52*, *p53*, *p21*, and *bcl-2*. The target agent can be an antibody that is present only in a subject infected with HIV-1, a viral protein antigen, or a protein characteristic of a disease state in a subject. The methods and apparatus of the present invention
 15 can be used for determining whether a subject is infected by a virus, whether nucleic acid obtained from a subject exhibits a single nucleotide mutation (SNM) relative to corresponding wild-type nucleic acid sequence, or whether a subject expresses a protein of interest, such as a bacterial protein, a fungal protein, a viral protein, an HIV protein, a hepatitis C protein, a hepatitis B protein, or a protein
 20 known to be specifically associated with a disease. An example of a dual bead experiment detecting a nucleic acid target is presented below in Example 1.

According to another aspect of the invention, there is provided multiplexing methods wherein more than one target agent (e.g., tens, hundreds, or even thousands of different target agents) can be identified on one optical analysis disc.
 25 Multiple capture agents can be provided in a single chamber together in capture fields, or separately in separate capture fields. Different reporter beads can be used to be distinguishable from each other, such as beads that fluoresce at different wavelengths or different size reporter beads. Experiments were performed to identify two different targets using the multiplexing technique. An example of one
 30 such assay is discussed in Example 2.

In accordance with yet another aspect, the invention includes an optical disc with a substrate, a capture layer associated with the substrate, and a capture agent bound to the capture layer, such that the capture agent binds to a dual bead complex. Multiple different capture agents can be used for different types of dual

bead complexes. The disc can be designed to allow for some dual bead processing on the disc with appropriate chambers and fluidic structures, and can be pre-loaded with reporter and capture beads so that only a sample needs to be added to form the dual bead complex structures.

5 According to still a further aspect of this invention, there is provided a disc and disc drive system for performing dual bead assays. The disc drive can include an electromagnet for performing the isolation process, and may include appropriate light source control and detection for the type of reporter beads used. The disc drive can be optical or magneto-optical.

10 For processing performed on the disc, the drive may advantageously include an electromagnet, and the disc preferably has a mixing chamber, a waste chamber, and capture area. In this embodiment, the sample is mixed with beads in the mixing chamber, a magnetic field is applied adjacent the mixing chamber, and the sample not held by the magnet is directed to the waste chamber so that all magnetic beads, 15 whether bound into a dual bead complex or unbound, remain in the mixing chamber. The magnetic beads are then directed to the capture area. One of a number of different valving arrangements can be used to control the flow. In still another aspect of the present invention, a bio-disc is produced for use with biological samples and is used in conjunction with a disc drive, such as a magneto-optical disc 20 drive, that can form magnetic regions on a disc. In a magneto-optical disc and drive, magnetic regions can be formed in a highly controllable and precise manner. These regions may be employed advantageously to magnetically bind magnetic beads, including unbound magnetic capture beads or including dual bead complexes with magnetic capture beads. The magneto-optical disc drive can write 25 to selected locations on the disc, and then use an optical reader to detect features located at those regions. The regions can be erased, thereby allowing the beads to be released.

In still another aspect, the invention includes a method for use with a bio-disc and drive including forming magnetic regions on the bio-disc, and providing 30 magnetic beads to the discs so that the beads bind at the magnetic locations. The method preferably further includes detecting at the locations where the magnetic beads bind biological samples, preferably using reporter beads that are detectable, such as by fluorescence or optical event detection. The method can be formed in multiple stages in terms of time or in terms of location through the use of multiple

chambers. The regions are written to and a sample is moved over the magnetic regions in order to capture magnetic beads. The regions can then be erased and released if desired. This method allows many different tests to be performed at one time, and can allow a level of interactivity between the user and the disc drives such

5 that additional tests can be created during the testing process.

It is also an aspect of the present invention to identify the key factors in optimizing the sensitivity of the dual bead assay. In a preferred embodiment thereof, methods have been brought out to select appropriate bead type that is, in turn, aimed at enhancing the covalent conjugation efficiency of DNA probe to beads.

10 Further, methods have been brought out to reduce the non-specific binding in the dual bead assays. In these methods, the capture and reporter beads are pretreated with various detergents prior to target capture. The bead pretreatments evaluated herein included anionic detergents (sodium dodecyl sulfate, dextran sulfate), cationic detergents (cetyl trimethylammonium bromide), non-ionic detergents

15 (CHAPS, octyl glucoside), phospholipids (phosphotidyl choline dipalmitoyl, phosphotidyl glycerol dipalmitoyl, phosphotidyl choline dilauroyl) and sheared denatured salmon sperm DNA. The results from several experiments on bead pre-treatment showed that the method most efficient in reducing the non-specific binding included pre-treating the capture and reporter beads with sheared denatured

20 salmon sperm DNA for one hour at room temperature prior to use in a dual bead assay. Pretreatment of capture and reporter beads with 10 μ g/ml denatured and sheared salmon sperm DNA reduced the non-specific binding approximately 10 fold. Data collected from one of the bead blocking experiments using detergents and salmon sperm DNA is presented in Fig. 38. To prevent any potential hybridization

25 between the salmon sperm DNA and the capture probes, the salmon sperm DNA was sheared prior to use.

Another aspect of the present invention is to arrive at methods for optimizing the ionic strength of the buffer used in the dual bead assays. The ionic strength of the assay medium plays a crucial factor in reducing the non-specific binding. The

30 non-specific binding of the dual bead assay was evaluated in the presence of a wide range of salt concentrations ranging from 0.1M up to 1.0M. It is a further aspect of the present invention to identify an optimized composition of the wash buffer since this plays a key role in reducing the non-specific binding. Figs. 40, 41, 42, and 43

illustrate results from various experiments related to buffer salt concentration titration and its effect on bead binding in a dual bead assay format.

In still another aspect, this invention is directed to methods for determining appropriate density of DNA probes on the solid phase. There exists a typical issue
5 in determining the desirable density of the probes. On one hand, a high concentration of the probes is required to obtain high conjugation efficiency. On the other hand, high density of the probes on the capture and reporter beads increases the non-specific binding and reduces the sensitivity of the dual bead assay. A high density of probes allows for interactions between the capture probes and reporter
10 probes in the absence of target, resulting in an increase in non-specific binding. The sensitivity of the dual bead assay would be optimal if only one target molecule binds to the capture beads and reporter beads. The number of reporter beads bound to the capture bead in the dual bead complex would then represent the number of target molecules. However, when the density of probes is high, at low target
15 concentrations, multiple target molecules can bind to the same capture bead. In this case, the number of reporter beads may no longer correspond to the number of target molecules.

The approach to this problem relies on reducing the density of the probes without compromising the probe conjugation efficiency. In this approach, the
20 capture and reporter beads are treated with probe blocking agents prior to target capture. The probe blocking agents may be DNA sequences complementary to the capture or reporter probes. They can bind to the capture or reporter probes and therefore significantly reduce the amount of available probes. The concentration of blocking agents are selected so that after incubation with the blocking agents only a
25 small fraction of capture or reporter probes are available. The use of probe blocking agents to increase sensitivity of an assay is not limited to beads but can be applied to any solid phase. An illustration of the use of a probe blocking agent in a bead type assay is presented in Fig. 44.

The formation of double stranded sequences between the blocking agents
30 and the capture or reporter probes makes the probes much more rigid, thus reducing the non-specific interactions between the capture beads and reporter beads. This results in a much lower non-specific binding in the dual bead assay.

Reducing the density of probes on the capture and reporter beads by using the blocking agents improves the sensitivity of the dual bead assay significantly by

increasing the ratio of signal (number of beads bound) to noise (non-specific binding).

In yet another aspect, the invention relates to identifying the desirable mode and time of mixing in a test method whereby the sensitivity of the dual bead assay is optimized. In a preferred embodiment, an efficient mixing method is brought out whereby the target mediated bead assay is improved several fold. Mixing of the capture beads during hybridization reaction is crucial. This is because if the magnetic beads remain stationary, very little target capture is observed. On the other hand, if the magnetic beads are mixed continuously, the target capture efficiency is significantly improved. According to one preferred embodiment of this aspect of the present invention, the most efficient mixing method included intermittent mixing, whereby the mixing is programmed so that the beads are mixed only when they start to settle down in the tube or on the disc. Experimental results from studies on mixing methods to increase dual bead formation are shown in Figs. 45 and 46.

The present invention also addresses implementing the methods recited above on to an analysis disc, modified optical disc, or an optical bio-disc. A bio-disc drive assembly may be employed to rotate the disc, read and process any encoded information stored on the disc, and analyze the DNA samples in the flow channel of the bio-disc. The bio-disc drive is thus provided with a motor for rotating the bio-disc, a controller for controlling the rate of rotation of the disc, a processor for processing return signals from the disc, and an analyzer for analyzing the processed signals. The rotation rate of the motor is controlled to achieve the desired rotation of the disc. The bio-disc drive assembly may also be utilized to write information to the bio-disc either before, during, or after the test material in the flow channel and target zones is interrogated by the read beam of the drive and analyzed by the analyzer. The bio-disc may include encoded information for controlling the rotation rate of the disc, providing processing information specific to the type of DNA test to be conducted, and for displaying the results on a monitor associated with the bio-drive.

The apparatus and methods of the different embodiments of the present invention can be designed for use by an end-user, inexpensively, without specialized expertise and expensive equipment. The system can be made portable, and thus usable in remote locations where traditional diagnostic equipment may not generally be available. Other related aspects applicable to components of this

assay system and signal acquisition methods are disclosed in commonly assigned and co-pending U.S. Patent Application Serial No. 10/038,297 entitled "Dual Bead Assays Including Covalent Linkages For Improved Specificity And Related Optical Analysis Discs" filed January 4, 2002; U.S. Provisional Application Serial No. 5 60/272,525 entitled "Biological Assays Using Dual Bead Multiplexing Including Optical Bio-Disc and Related Methods" filed March 1, 2001; and U.S. Provisional Application Serial Nos. 60/275,643, 60/314,906, and 60/352,270 each entitled "Surface Assembly for Immobilizing Capture Agents and Dual Bead Assays Including Optical Bio-Disc and Methods Relating Thereto" respectively filed March 10 14, 2001, August 24, 2001, and January 30, 2002. All of these applications are herein incorporated by reference in their entirety.

Other features and advantages of the different embodiments and aspects of the present invention will become apparent from the following detailed description, accompanying drawing figures, and related examples.

BRIEF DESCRIPTION OF THE DRAWING FIGURES

Further objects of the present invention together with additional features contributing thereto and advantages accruing therefrom will be apparent from the following description of preferred embodiments of the present invention which are 20 shown in the accompanying drawing figures with like reference numerals indicating like components throughout, wherein:

Fig. 1 is a perspective view of an optical disc system according to the present invention;

Fig. 2 is a block and pictorial diagram of an optical reading system according 25 to embodiments of the present invention;

Figs. 3A, 3B, and 3C are respective exploded, top, and perspective views of a reflective disc according to embodiments of the present invention;

Figs. 4A, 4B, and 4C are respective exploded, top, and perspective views of a transmissive disc according to embodiments of the present invention;

Fig. 5A is a partial longitudinal cross sectional view of the reflective optical bio-disc shown in Figs. 3A, 3B, and 3C illustrating a wobble groove formed therein; 30

Fig. 5B is a partial longitudinal cross sectional view of the transmissive optical bio-disc illustrated in Figs. 4A, 4B, and 4C showing a wobble groove formed therein and a top detector;

Fig. 6A is a partial radial cross-sectional view of the disc illustrated in Fig. 5A;

Fig. 6B is a partial radial cross-sectional view of the disc illustrated in Fig. 5B;

5 Figs. 7A, 8A, 9A, and 10A are schematic representations of a capture bead, a reporter bead, and a dual bead complex as utilized in conjunction with genetic assays;

Figs. 7B, 8B, 9B, and 10B are schematic representations of a capture bead, a reporter bead, and a dual bead complex as employed in conjunction with immunochemical assays;

10 Fig. 11A is a pictorial representation of one embodiment of a method for producing genetic dual bead complex solutions;

Fig. 11B is a pictorial representation of one embodiment of a method for producing immunochemical dual bead complex solutions;

Fig. 12A is a pictorial representation of another embodiment of a method for producing genetic dual bead complex solutions;

15 Fig. 12B is a pictorial representation of another embodiment of a method for producing immunochemical dual bead complex solutions;

Fig. 13 is a longitudinal cross sectional view illustrating the disk layers in combination with a mixing or loading chamber;

20 Fig. 14 is a view similar to Fig. 13 showing the mixing chamber loaded with dual bead complex solution;

Figs. 15A and 15B are radial cross sectional views of the disc and target zone illustrating one embodiment for binding of reporter beads to capture agents in a genetic assay;

25 Figs. 16A and 16B are radial cross sectional views of the disc and target zone showing another embodiment for binding of reporter beads to capture agents in a genetic assay;

Fig. 17 is radial cross sectional view of the disc and target zone illustrating one embodiment for binding of capture beads to capture agents in a genetic assay;

30 Fig. 18 is radial cross sectional view of the disc and target zone depicting another embodiment for binding of capture beads to capture agents in a genetic assay;

Figs. 19A, 19B, and 19C are partial cross sectional views illustrating one embodiment of a method according to this invention for binding the reporter bead of a dual bead complex to a capture layer in a genetic assay;

Figs. 20A, 20B, and 20C are partial cross sectional views showing one embodiment of a method according to the present invention for binding the reporter bead of a dual bead complex to a capture layer in an immunochemical assay;

5 Figs. 21A, 21B, and 21C are partial cross sectional views illustrating another embodiment of a method according to this invention for binding the reporter bead of a dual bead complex to a capture layer in a genetic assay;

Figs. 22A, 22B, and 22C are partial cross sectional views presenting another embodiment of a method according to the invention for binding the reporter bead of a dual bead complex to a capture layer in an immunochemical assay;

10 Figs. 23A and 23B are partial cross sectional views depicting one embodiment of a method according to the present invention for binding the capture bead of a dual bead complex to a capture layer in a genetic assay;

Figs. 24A and 24B are partial cross sectional views showing another embodiment of a method according to this invention for binding the capture bead of a dual bead complex to a capture layer in a genetic assay;

Figs. 25A–25D illustrate a method according to the present invention for detecting the presence of target DNA or RNA in a genetic sample utilizing an optical bio-disc;

20 Figs. 26A–26D illustrate another method according to this invention for detecting the presence of target DNA or RNA in a genetic sample utilizing an optical bio-disc;

Figs. 27A–27D illustrate a method according to the present invention for detecting the presence of a target antigen in a biological test sample utilizing an optical bio-disc;

25 Fig. 28A is a graphical representation of an individual 2.1 micron reporter bead and a 3 micron capture bead positioned relative to the tracks of an optical bio-disc according to the present invention;

Fig. 28B is a series of signature traces derived from the beads of Fig. 28A utilizing a detected signal from the optical drive according to the present invention;

30 Fig. 29A is a graphical representation of a 2.1 micron reporter bead and a 3 micron capture bead linked together in a dual bead complex positioned relative to the tracks of an optical bio-disc according to the present invention;

Fig. 29B is a series of signature traces derived from the dual bead complex of Fig. 29A utilizing a detected signal from the optical drive according to this invention;

Fig. 30A is a bar graph showing results from a dual bead assay according to the present invention;

Fig. 30B is a graph showing a standard curve demonstrating the detection limit for fluorescent beads detected with a fluorimeter;

5 Fig. 30C is a pictorial representation demonstrating the formation of the dual bead complex;

Fig. 31 is a bar graph showing the sensitivity of disc drive detection using a dual bead complex;

10 Fig. 32 is a schematic representation of combining beads for dual bead assay multiplexing according to embodiments of the present invention;

Fig. 33A is a schematic representation of a fluidic circuit according to the present invention utilized in conjunction with a magnetic field generator to control movement of magnetic beads;

15 Figs. 33B-33D are schematics of a first fluidic circuit that implements the valving structure of FIG. 33A according to one embodiment of fluid transport aspects of the present invention;

Figs. 34A-34C are schematics of a second fluidic circuit that implements the valving structure of FIG. 33A according to another embodiment of fluid transport aspects of the present invention;

20 Fig. 35 is a perspective view of a the magnetic field generator and a disc including one embodiment of a fluidic circuit employed in conjunction with magnetic beads according to this invention;

Figs. 36A, 36B, and 36C are plan views illustrating a method of separation and detection for dual bead assays using the fluidic circuit shown in Fig. 35;

25 Fig. 37 is a perspective view of a magneto-optical bio-disc showing magnetic regions, magnetically bound capture beads, and the formation of dual bead complexes according to another aspect of the present invention;

Fig. 38 is a bar graph presentation demonstrating the pretreatment of the beads with various blocking agents indicating that use of salmon sperm DNA
30 worked best in reducing the nonspecific bead binding;

Fig.39 is a bar graph presentation showing the range of detection of the dual bead assay;

Fig. 40 is a bar graph illustrating the use of NaCl in varying concentrations and the related non-specific binding;

Fig. 41 is a bar graph presentation showing increasing EDTA concentration and the related non-specific binding;

Fig. 42 is a bar graph presentation depicting an increasing NaCl concentration and the related non-specific binding;

5 Figs. 43A and 43B are bar graph presentations illustrating an increasing concentration of $MgCl_2$ and related non-specific binding;

Fig. 44 is a pictorial schematic representation showing the use of probe blocking agents to increase the sensitivity of the bead assay;

Fig. 45 is a bar graph presentation illustrating the effect of incubation time
10 during a hybridization reaction; and

Fig. 46 is a bar graph presentation showing a mixing method directed to increasing efficiency in dual bead binding.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

15 The following description of the present invention relates to optical analysis discs, disc drive systems, and assay chemistries and techniques. The invention further relates to alternate magneto-optical drive systems, MO bio-discs, and related processing methods.

20 Disc Drive System and Related Optical Analysis discs

With reference now to Fig. 1, there is shown a perspective view of an optical bio-disc 110 for use in an optical disc drive 112. Drive 112, in conjunction with software in the drive or associated with a separate computer, can cause images, graphs, or output data to be displayed on display monitor 114. As indicated below,
25 there are different types of discs and drives that can be used. The disc drive can be in a unit separate from a controlling computer, or provided in a bay within a computer. The device can be made as portable as a laptop computer, and thus usable with battery power and in remote locations not generally served by advanced diagnostic equipment. The drive is preferably a conventional drive with minimal or
30 no hardware modification, but can be a dedicated bio-disc drive. Further details regarding these types of drive systems and related signal processing methods are disclosed in, for example, commonly assigned and co-pending U.S. Patent Application Serial No. 09/378,878 entitled "Methods and Apparatus for Analyzing Operational and Non-operational Data Acquired from Optical Discs" filed August 23,

1999; U.S. Provisional Patent Application Serial No. 60/150,288 entitled "Methods and Apparatus for Optical Disc Data Acquisition Using Physical Synchronization Markers" filed August 23, 1999; U.S. Patent Application Serial No. 09/421,870 entitled "Trackable Optical Discs with Concurrently Readable Analyte Material" filed
 5 October 26, 1999; U.S. Patent Application Serial No. 09/643,106 entitled "Methods and Apparatus for Optical Disc Data Acquisition Using Physical Synchronization Markers" filed August 21, 2000; U.S. and U.S. Patent Application Serial No. 10/043,688 entitled "Optical Disc Analysis System Including Related Methods For Biological and Medical Imaging" filed January 10, 2002. These applications are
 10 herein incorporated by reference in their entirety.

Optical bio-disc 110 for use with embodiments of the present invention may have any suitable shape, diameter, or thickness, but preferably is implemented on a round disc with a diameter and a thickness similar to those of a compact disc (CD), a recordable CD (CD-R), CD-RW, a digital versatile disc (DVD), DVD-R, DVD-RW,
 15 or other standard optical disc format. The disc may include encoded information, preferably in a known format, for performing, controlling, and post-processing a test or assay, such as information for controlling the rotation rate and direction of the disc, timing for rotation, stopping and starting, delay periods, locations of samples, position of the light source, and power of the light source. Such encoded
 20 information is referred to generally here as operational information.

The disc may be a reflective disc, as shown in Figs. 3A-3C, a transmissive disc, Figs. 4A-4C, or some combination of reflective and transmissive. In a reflective disc, an incident light beam is focused onto the disc (typically at a reflective surface where information is encoded), reflected, and returned through
 25 optical elements to a detector on the same side of the disc as the light source. In a transmissive disc, light passes through the disc (or portions thereof) to a detector on the other side of the disc from the light source. In a transmissive portion of a disc, some light may also be reflected and detected as reflected light.

Fig. 2 shows an optical disc reader system 116. This system may be a
 30 conventional reader for CD, CD-R, DVD, or other known comparable format, a modified version of such a drive, or a completely distinct dedicated device. The basic components are a motor for rotating the disc, a light system for providing light, and a detection system for detecting light.

With reference now generally to Figs. 2 to 4C, a light source 118 provides light to optical components 120 to produce an incident light beam 122. In the case of reflective disc 144, Figs. 3A-3C, a return beam 124 is reflected from either reflective surface 156, 174, or 186, Figs. 3C and 4C. Return beam 124 is provided
 5 back to optical components 120, and then to a bottom detector 126. In this type of disc, the return beam may carry operational information or other encoded data as well as characteristic information about the investigational feature or test sample under study.

For transmissive disc 180, Figs. 4A-4C, some of the energy from the incident
 10 beam 122 will undergo a light/matter interaction with an investigational feature or test sample and then proceed through the disc as a transmitted beam 128 that is detected by a top detector 130. For a transmissive disc including a semi-reflective layer 186 (Fig. 4C) as the operational layer, some of the energy from the incident beam 122 will also reflect from the operational layer as return beam 124 which
 15 carries operational information or stored data. Optical components 120 can include a lens, a beam splitter, and a quarter wave plate that changes the polarization of the light beam so that the beam splitter directs a reflected beam through the lens to focus the reflected beam onto the detector. An astigmatic element, such as a cylindrical lens, may be provided between the beam splitter and detector to
 20 introduce astigmatism in the reflected light beam. The light source can be controllable to provide variable wavelengths and power levels over a desired range in response to data introduced by the user or read from the disc. This controllability is especially useful when it is desired to detect multiple different structures that fluoresce at different wavelengths.

25 Now with continuing reference to Fig. 2, it is shown that data from detector 126 and/or detector 130 is provided to a computer 132 including a processor 134 and an analyzer 136. An image or output results can then be provided to a monitor 114. Computer 132 can represent a desktop computer, programmable logic, or some other processing device, and also can include a connection (such as over the
 30 Internet) to other processing and/or storage devices. A drive motor 140 and a controller 142 are provided for controlling the rotation rate and direction or rotation of disc the 144 or 180. Controller 142 and the computer 132 with processor 134 can be in remote communication or implemented in the same computer. Methods and

systems for reading such a disc are also shown in Gordon, U.S. Patent No. 5,892,577, which is incorporated herein by reference.

The detector can be designed to detect all light that reaches the detector, or though its design or an external filter, light only at specific wavelengths. By making
5 the detector controllable in terms of the detectable wavelength, beads or other structures that fluoresce at different wavelengths can be separately detected.

A hardware trigger sensor 138 may be used with either a reflective disc 144 or transmissive disc 180. Triggering sensor 138 provides a signal to computer 132 (or to some other electronics) to allow for the collection of data by processor 134
10 only when incident beam 122 is on a target zone or inspection area. Alternatively, software read from a disc can be used to control data collection by processor 134 independent of any physical marks on the disc. Such software or logical triggering is discussed in further detail in commonly assigned and co-pending U.S. Provisional Application Serial No. 60/352,625 entitled "Logical Triggering Methods And
15 Apparatus For Use With Optical Analysis Discs And Related Disc Drive Systems" filed January 28, 2002, which is herein incorporated by reference in its entirety.

The substrate layer of the optical analysis disc may be impressed with a spiral track that starts at an innermost readable portion of the disc and then spirals out to an outermost readable portion of the disc. In a non-recordable CD, this track
20 is made up of a series of embossed pits with varying length, each typically having a depth of approximately one-quarter the wavelength of the light that is used to read the disc. The varying lengths and spacing between the pits encode the operational data. The spiral groove of a recordable CD-like disc has a detectable dye rather than pits. This is where the operation information, such as the rotation rate, is
25 recorded. Depending on the test, assay, or investigational protocol, the rotation rate may be variable with intervening or consecutive periods of acceleration, constant speed, and deceleration. These periods may be closely controlled both as to speed and time of rotation to provide, for example, mixing, agitation, or separation of fluids and suspensions with agents, reagents, antibodies, or other materials. Different
30 optical analysis disc and bio-disc designs that may be utilized with the present invention, or readily adapted thereto, are disclosed, for example, in commonly assigned, copending U.S. Patent Application Serial No. 09/999,274 entitled "Optical Bio-discs with Reflective Layers" filed on November 15, 2001; U.S. Patent Application Serial No. 10/005,313 entitled "Optical Discs for Measuring Analytes"

filed December 7, 2001; U.S. Patent Application Serial No. 10/006,371 entitled "Methods for Detecting Analytes Using Optical Discs and Optical Disc Readers" filed December 10, 2001; U.S. Patent Application Serial No. 10/006,620 entitled "Multiple Data Layer Optical Discs for Detecting Analytes" filed December 10, 2001; and U.S. Patent Application Serial No. 10/006,619 entitled "Optical Disc Assemblies for Performing Assays" filed December 10, 2001, which are all herein incorporated by reference in their entirety.

Numerous designs and configurations of an optical pickup and associated electronics may be used in the context of the embodiments of the present invention.

Further details and alternative designs for compact discs and readers are described in *Compact Disc Technology*, by Nakajima and Ogawa, IOS Press, Inc. (1992); *The Compact Disc Handbook, Digital Audio and Compact Disc Technology*, by Baert et al. (eds.), Books Britain (1995); and *CD-Rom Professional's CD-Recordable Handbook: The Complete Guide to Practical Desktop CD*, Starrett et al. (eds.), ISBN:0910965188 (1996); all of which are incorporated herein in their entirety by reference.

The disc drive assembly is thus employed to rotate the disc, read and process any encoded operational information stored on the disc, and analyze the liquid, chemical, biological, or biochemical investigational features in an assay region of the disc. The disc drive assembly may be further utilized to write information to the disc either before, during, or after the material in the assay zone is analyzed by the read beam of the drive. In alternate embodiments, the disc drive assembly is implemented to deliver assay information through various possible interfaces such as via Ethernet to a user, over the Internet, to remote databases, or anywhere such information could be advantageously utilized. Further details relating to this type of disc drive interfacing are disclosed in commonly assigned copending U.S. Patent Application Serial No. 09/986,078 entitled "Interactive System For Analyzing Biological Samples And Processing Related Information And The Use Thereof " filed November 7, 2001, which is incorporated herein by reference in its entirety.

Referring now specifically to Figs. 3A, 3B, and 3C, the reflective disc 144 is shown with a cap 146, a channel layer 148, and a substrate 150. The channel layer 148 may be formed by a thin-film adhesive member. Cap 146 has inlet ports 152 for receiving samples and vent ports 154. Cap 146 may be formed primarily from

polycarbonate, and may be coated with a cap reflective layer 156 on the bottom thereof. Reflective layer 156 is preferably made from a metal such as aluminum or gold.

Channel layer 148 defines fluidic circuits 158 by having desired shapes cut out from channel layer 148. Each fluidic circuit 158 preferably has a flow channel 160 and a return channel 162, and some have a mixing chamber 164. A mixing chamber 166 can be symmetrically formed relative to the flow channel 160, while an off-set mixing chamber 168 is formed to one side of the flow channel 160. Fluidic circuits 158 are rather simple in construction, but a fluidic circuit can include other channels and chambers, such as preparatory regions or a waste region, as shown, for example, in U.S. Patent No. 6,030,581, which is incorporated herein by reference, and can include valves and other fluid control structures. Channel layer 148 can include adhesives for bonding to the substrate and to the cap.

Substrate 150 has a plastic layer 172, and has target zones 170 formed as openings in a substrate reflective layer 174 deposited on the top of layer 172. In this embodiment, reflective layer 174, best illustrated in Fig. 3C, is used to encode operational information. Plastic layer 172 is preferably formed from polycarbonate. Target zones 170 may be formed by removing portions of the substrate reflective layer 174 in any desired shape, or by masking target zone areas before applying substrate reflective layer 174. The substrate reflective layer 174 is preferably formed from a metal, such as aluminum or gold, and can be configured with the rest of the substrate to encode operational information that is read with incident light, such as through a wobble groove or through an arrangement of pits. Light incident from under substrate 150 thus is reflected by layer 174, except at target zones 170, where it is reflected by layer 156. Target zones are where investigational features are detected. If the target zone is a location where an antibody, strand of DNA, or other material that can bind to a target is located, the target zone can be referred to as a capture zone.

With reference now particularly to Fig. 3C, optical disc 144 is cut away to illustrate a partial cross-sectional perspective view. An active layer 176 is formed over substrate reflective layer 174. Active layer 176 may generally be formed from nitrocellulose, polystyrene, polycarbonate, gold, activated glass, modified glass, or a modified polystyrene such as, for example, polystyrene-co-maleic anhydride. In this embodiment, channel layer 148 is situated over active layer 174.

In operation, samples can be introduced through inlet ports 152 of cap 146. When rotated, the sample moves outwardly from inlet port 152 along active layer 176. Through one of a number of biological or chemical reactions or processes, detectable features, referred to as investigational features, may be present in the target zones. Examples of such processes are shown in the incorporated U.S. Patent No. 6,030,581 and commonly assigned, co-pending U.S. Patent Application No. 09/988,728 entitled "Methods And Apparatus For Detecting And Quantifying Lymphocytes With Optical Biodiscs" filed November 16, 2001; and U.S. Patent Application No. 10/035,836 entitled "Surface Assembly For Immobilizing DNA Capture Probes And Bead-Based Assay Including Optical Bio-Discs And Methods Relating Thereto" filed December 21, 2001, both of which are herein incorporated by reference in their entireties.

The investigational features captured within the target zones, by the capture layer with a capture agent, may be designed to be located in the focal plane coplanar with reflective layer 174, where an incident beam is typically focused in conventional readers. Alternatively, the investigational features may be captured in a plane spaced away from the focal plane. The former configuration is referred to as a "proximal" type disc, and the latter a "distal" type disc.

Referring to Figs. 4A, 4B, and 4C, it is shown that one particular embodiment of the transmissive optical disc 180 includes a clear cap 182, a channel layer 148, and a substrate 150. The clear cap 182 includes inlet ports 152 and vent ports 154 and is preferably formed mainly from polycarbonate. Trigger marks 184 may be included on the cap 182. Channel layer 148 has fluidic circuits 158, which can have structure and use similar to those described in conjunction with Figs. 3A, 3B, and 3C. Substrate 150 may include target zones 170, and preferably includes a polycarbonate layer 172. Substrate 150 may, but need not, have a thin semi-reflective layer 186 deposited on top of layer 172. Semi-reflective layer 186 is preferably significantly thinner than substrate reflective layer 174 on substrate 150 of reflective disc 144 (Figs. 3A-3C). Semi-reflective layer 186 is preferably formed from a metal, such as aluminum or gold, but is sufficiently thin to allow a portion of an incident light beam to penetrate and pass through layer 186, while some of the incident light is reflected back. A gold film layer, for example, is 95% reflective at a thickness greater than about 700 Å, while the transmission of light through the gold film is about 50% transmissive at approximately 100 Å.

Fig. 4C is a cut-away perspective view of transmissive disc 180. The semi-reflective nature of layer 186 makes its entire surface potentially available for target zones, including virtual zones defined by trigger marks or encoded data patterns on the disc. Target zones 170 may also be formed by marking the designated area in the indicated shape or alternatively in any desired shape. Markings to indicate target zone 170 may be made on semi-reflective layer 186 or on a bottom portion of substrate 150 (under the disc). Target zones 170 may be created by silk screening ink onto semi-reflective layer 186.

An active layer 176 is applied over semi-reflective layer 186. Active layer 176 may be formed from the same materials as described above in conjunction with layer 176 (Fig. 3C) and serves substantially the same purpose when a sample is provided through an opening in disc 180 and the disc is rotated. In transmissive disc 180, there is no reflective layer, on the clear cap 182, comparable to reflective layer 156 in reflective disc 144 (Fig. 3C).

Referring now to Fig. 5A, there is shown a cross sectional view taken across the tracks of the reflective disc embodiment 144 (Figs. 3A-3C) of the bio-disc 110 (Fig. 1) according to the present invention. As illustrated, this view is taken longitudinally along a radius and flow channel of the disc. Fig. 5A includes the substrate 150 which is composed of a plastic layer 172 and a substrate reflective layer 174. In this embodiment, the substrate 150 includes a series of grooves 188. The grooves 188 are in the form of a spiral extending from near the center of the disc toward the outer edge. The grooves 188 are implemented so that the interrogation or incident beam 122 may track along the spiral grooves 188 on the disc. This type of groove 188 is known as a "wobble groove". The groove 188 is formed by a bottom portion having undulating or wavy side walls. A raised or elevated portion separates adjacent grooves 188 in the spiral. The reflective layer 174 applied over the grooves 188 in this embodiment is, as illustrated, conformal in nature. Fig. 5A also shows the active layer 176 applied over the reflective layer 174. As shown in Fig. 5A, the target zone 170 is formed by removing an area or portion of the reflective layer 174 at a desired location or, alternatively, by masking the desired area prior to applying the reflective layer 174. As further illustrated in Fig. 5A, the plastic adhesive member or channel layer 148 is applied over the active layer 176. Fig. 5A also shows the cap portion 146 and the reflective surface 156 associated therewith. Thus, when the cap portion 146 is applied to the plastic

adhesive member 148 including the desired cut-out shapes, the flow channel 160 is thereby formed.

Fig. 5B is a cross sectional view, similar to that illustrated in Fig. 5A, taken across the tracks of the transmissive disc embodiment 180 (Figs. 4A-4C) of the bio-disc 110 (Fig. 1) according to the present invention. This view is taken longitudinally along a radius and flow channel of the disc. Fig. 5B illustrates the substrate 150 that includes the thin semi-reflective layer 186. This thin semi-reflective layer 186 allows the incident or interrogation beam 122, from the light source 118 (Fig. 2), to penetrate and pass through the disc to be detected by the top detector 130, while some of the light is reflected back in the form of the return beam 124. The thickness of the thin semi-reflective layer 186 is determined by the minimum amount of reflected light required by the disc reader to maintain its tracking ability. The substrate 150 in this embodiment, like that discussed in Fig. 5A, includes the series of grooves 188. The grooves 188 in this embodiment are also preferably in the form of a spiral extending from near the center of the disc toward the outer edge. The grooves 188 are implemented so that the interrogation beam 122 may track along the spiral. Fig. 5B also shows the active layer 176 applied over the thin semi-reflective layer 186. As further illustrated in Fig. 5B, the plastic adhesive member or channel layer 148 is applied over the active layer 176. Fig. 5B also shows the clear cap 182. Thus, when the clear cap 182 is applied to the plastic adhesive member 148 including the desired cut-out shapes, the flow channel 160 is thereby formed and a part of the incident beam 122 is allowed to pass therethrough substantially unreflected. The amount of light that passes through can then be detected by the top detector 130.

Fig. 6A is a view similar to Fig. 5A but taken perpendicularly to a radius of the disc to illustrate the reflective disc and the initial refractive property thereof when observing the flow channel 160 from a radial perspective. In a parallel comparison manner, Fig. 6B is a similar view to Fig. 5B but taken perpendicularly to a radius of the disc to represent the transmissive disc and the initial refractive property thereof when observing the flow channel 160 from a radial perspective. Grooves 188 are not seen in Figs. 5A and 5B since the sections are cut along the grooves 188. Figs. 6A and 6B show the presence of the narrow flow channel 160 that is situated perpendicular to the grooves 188 in these embodiments. Figs. 5A, 5B, 6A, and 6B show the entire thickness of the respective reflective and transmissive discs. In

these views, the incident beam 122 is illustrated initially interacting with the substrate 150 which has refractive properties that change the path of the incident beam as shown to provide focusing of the beam 122 on the reflective layer 174 or the thin semi-reflective layer 186.

5

Assay Chemistries and Dual Bead Formation

Referring now to Figs. 7A-10A and 7B-10B, there is shown a capture bead 190, a reporter bead 192, and the formation of a dual bead complex 194. Capture bead 190 can be used in conjunction with a variety of different assays including biological assays such as immunoassays (Figs. 7B-10B), molecular assays, and more specifically genetic assays (Figs. 7A-10A). In the case of immunoassays, antibody transport probes 196 are conjugated onto the beads. Antibody transport probes 196 include proteins, such as antigens or antibodies, implemented to capture protein targets. In the case of molecular assays, oligonucleotide transport probes 198 would be conjugated onto the beads. Oligonucleotide transport probes 198 include nucleic acids such as DNA or RNA implemented to capture genetic targets.

As shown in Fig. 7A, a target agent such as target DNA or RNA 202, obtained from a test sample, is added to a capture bead 190 coated with oligonucleotide transport probes 198. In this implementation, transport probes 198 are formed from desired sequences of nucleic acids. Related aspects directed to DNA probe conjugation onto solid phase of this system of assays are discussed in further detail in commonly assigned and co-pending U.S. Provisional Application Serial No. 60/278,685 entitled "Use of Double Stranded DNA for Attachment to Solid Phase to Reduce Non-Covalent Binding" filed March 26, 2001. This application is herein incorporated by reference in its entirety.

As shown in Fig. 7B, a target agent such as target antigen 204 from a test sample is added to a capture bead 190 coated with antibody transport probes 196. In this alternate implementation, the transport probes 196 are formed from proteins such as antibodies.

Capture bead 190 has a characteristic that allows it to be isolated from a material suspension or solution. The capture bead may be selected based upon a desired size, and a preferred way to make it isolatable is for it to be magnetic.

Fig. 8A illustrates the binding of target DNA or RNA 202 to complementary transport probes 198 on capture bead 190 in the genetic assay implementation of the present invention. Fig. 8B shows an immunoassay version of Fig. 8A, transport probes 196 can alternatively include antibodies or antigens for binding to a target protein 204.

Fig. 9A shows a reporter bead 192 coated with oligonucleotide signal probes 206 complementary to target agent 202 (see Fig. 8A). Reporter bead 192 is selected based upon a desired size and the material properties for detection and reporting purposes. In one specific embodiment a 2.1 micron polystyrene bead is employed. Signal probes 206 can be strands of DNA or RNA to capture target DNA or RNA.

Fig. 9B illustrates a reporter bead 192 coated with antibody signal probes 208 that bind to the target agent 204 as shown in Fig. 8B. Reporter bead 192 is selected based upon a desired size and the material properties for detection and reporting purposes. This may also preferably include a 2.1 micron polystyrene bead. Signal probes 208 can be antigens or antibodies implemented to capture protein or glycoprotein targets.

Fig. 10A is a pictorial representation of a dual bead complex 194 that can be formed from capture bead 190 with probe 198, target agent 202, and reporter bead 192 with probe 206. Probes 198 and 206 conjugated on capture bead 190 and reporter bead 192, respectively, have sequences complementary to the target agent 202, but not to each other. Further details regarding target agent detection and methods of reducing non-specific binding of target agents to beads are discussed in commonly assigned and co-pending U.S. Provisional Application Serial No. 60/278,106 entitled "Dual Bead Assays Including Use of Restriction Enzymes to Reduce Non-Specific Binding" filed March 23, 2001; and U.S. Provisional Application Serial No. 60/278,110 entitled "Dual Bead Assays Including Use of Chemical Methods to Reduce Non-Specific Binding" filed March 23, 2001, which are both incorporated herein by reference in their entirety.

Fig. 10B is a pictorial representation of the immunoassay version of a dual bead complex 194 that can be formed from capture bead 190 with probe 196, target agent 204, and reporter bead 192 with probe 208. Probes 196 and 208 conjugated on capture bead 190 and reporter bead 192, respectively, only bind to the target agent 202, and not to each other.

In an alternative embodiment of the current system of assays, target agent binding efficiency and specificity may be enhanced by using a cleavable spacer that temporarily links the reporter bead 192 and capture bead 190. The dual bead complex formed by the cleavable spacer essentially places the transport probe and the signal probe in close proximity to each other thus allowing more efficient target binding to both probes. Once the target agent is bound to the probes the spacer may then be cleaved permitting the bound target agent to retain the dual bead structure. The use of cleavable spacers in dual bead assay systems is disclosed in further detail in commonly assigned and co-pending U.S. Provisional Application Serial No. 60/278,688 entitled "Dual Bead Assays Using Cleavable Spacers to Improve Specificity and Sensitivity" filed March 26, 2001, which by reference hereto is incorporated into this application in its entirety.

With reference now to Fig. 11A, there is illustrated a method of preparing a molecular assay using a "single-step hybridization" technique to create dual bead complex structures in a solution according to one aspect of the present invention. This method includes 5 principal steps identified consecutively as Steps I, II, III, IV, and V discussed in detail below. Generally, this particular method includes the steps of providing a mixture of capture beads that have transport probes covalently bound thereto, providing a mixture of reporter beads that have signal probes covalently bound thereto, blocking the mixture of capture beads with a bead blocking agent, blocking the mixture of reporter beads with the bead blocking agent, suspending the mixture of capture beads in a hybridization solution, adding to the mixture a target agent that hybridizes with the transport probes, adding to the mixture the reporter beads, allowing the signal probes to hybridize with the target agent to thereby form a dual bead complex including at least one capture bead and one reporter bead, separating the dual bead complex from unbound reporter beads, removing from the mixture the unbound reporter beads, and loading the mixture including the dual bead complex into an optical bio-disc for analysis.

More particularly now, in Step I of this method a number of capture beads 190 coated with oligonucleotide transport probes 198 are deposited into a test tube 212 containing a buffer solution 210. In this embodiment, the buffer solution 210 includes a blocking agent such as salmon sperm DNA, Denhart's solution, and BSA. The preferred blocking agent is salmon sperm DNA. The number of capture beads 190 used in this method may be, for example, on the order of $10E+07$ and each on

the order of 1 micron or greater in diameter. Capture beads 190 are suspended in hybridization solution and are loaded into the test tube 212 by injection with pipette 214. The preferred hybridization solution is composed of 0.2M NaCl, 10mM MgCl₂, 1mM EDTA, 50mM Tris-HCl, pH 7.5, and 5X Denhart's mix. A desirable
 5 hybridization temperature is 37 degrees Celsius. In a preliminary step in this embodiment, transport probes 198 are conjugated to 3 micron magnetic capture beads 190 by EDC conjugation. Further details regarding conjugation methods are disclosed in commonly assigned U.S. Provisional Application Serial No. 60/271,922 entitled, "Methods for Attaching Capture DNA and Reporter DNA to Solid Phase
 10 Including Selection of Bead Types as Solid Phase" filed February 27, 2001; and U.S. Provisional Application Serial No. 60/277,854 entitled "Methods of Conjugation for Attaching Capture DNA and Reporter DNA to Solid Phase" filed March 22, 2001, both of which are herein incorporated by reference in their entirety.

As shown in Step II, target DNA or RNA 202 is added to the solution.
 15 Oligonucleotide transport probes 198 are complementary to the DNA or RNA target agent 202. The target DNA or RNA 202 thus binds to the complementary sequences of transport probe 198 attached to the capture bead 190 as shown in Fig. 8A.

With reference now to Step III, there is added to the solution 210 reporter
 20 beads 192 coated with oligonucleotide signal probes 206. As also shown in Figs. 9A and 10A, signal probes 206 are complementary to the target DNA or RNA 202. In one embodiment, signal probes 206, which are complementary to a portion of the target DNA or RNA 202, are conjugated to 2.1 micron fluorescent reporter beads 192. Signal probes 206 and transport probes 198 each have sequences that are
 25 complementary to the target DNA 202, but not complementary to each other. After adding reporter beads 192, the dual bead complex 194 is formed such that the target DNA 202 links capture bead 190 and reporter beads 192. With specific and thorough washing, there should be minimal non-specific binding between reporter bead 192 and capture bead 190. The target agent 202 and signal probe 206 are
 30 preferably allowed to hybridize for three to four hours at 37 degrees Celsius.

In this embodiment and others, it was found that intermittent mixing (i.e., periodically mixing and then stopping) produced greater yield of dual bead complex than continuous mixing during hybridization.

As next shown in Step IV, after hybridization, the dual bead complex 194 is separated from unbound reporter beads in the solution. The solution can be exposed to a magnetic field to capture the dual bead complex structures 194 using the magnetic properties of capture bead 190. The magnetic field can be encapsulated in a magnetic test tube rack 216 with a built-in magnet 218, which can be permanent or electromagnetic to draw out the magnetic beads and remove any unbound reporter beads in the suspension. Note that capture beads not bound to reporter beads will also be isolated.

The purification process illustrated in Step IV includes the removal of supernatant containing free-floating particles. Wash buffer is added into the test tube and the bead solution is mixed well. The preferred wash buffer for the one step assay consists of 145mM NaCl, 50mM Tris, pH 7.5, 0.1% SDS, 0.05% Tween, 0.25% NFDM, and 10mM EDTA. Most of the unbound reporter beads 182, free-floating DNA, and non-specifically bound particles are agitated and removed from the supernatant. The dual bead complex can form a matrix of capture beads, target sequences, and reporter beads, wherein the wash process can further assist in the extraction of free floating particles trapped in the lattice structure of overlapping dual bead particles. Further details relating to other aspects associated with methods of decreasing non-specific binding of reporter beads to capture beads are disclosed in, for example, commonly assigned U.S. Provisional Application Serial No. 60/272,134 entitled "Reduction of Non-Specific Binding in Dual Bead Assays by Selection of Bead Type and Bead Treatment" filed February 28, 2001; and U.S. Provisional Application Serial No. 60/275,006 entitled "Reduction of Non-Specific Binding in Dual Bead Assays by Selection of Buffer Conditions and Wash Conditions" filed March 12, 2001. Both of these applications are herein incorporated by reference in their entirety.

The last principal step shown in Fig. 11A is Step V. In this step, once the dual bead complex has been washed approximately 3-5 times with wash buffer solution, the assay mixture may be loaded into the disc and ready to be analyzed.

Fig. 11B illustrates an immunoassay using a "single-step antigen binding" method, similar to that in Fig. 11A, to create dual bead complex structures in a solution. This method similarly includes 5 principal steps. These steps are respectively identified as Steps I, II, III, IV, and V in Fig. 11A.

As shown in Step I, capture beads 190, e.g., on the order of $10E+07$ in number and each on the order of 1 micron or above in diameter, which are coated with antibody transport probes 196 are added to a buffer solution 210. In this embodiment, the buffer solution 210 includes a blocking agent such as salmon sperm DNA, Denhart's solution, and BSA. The preferred blocking agent is BSA. This solution may be that same as that employed in the method shown in Fig. 11A or alternatively may be specifically prepared for use with immunochemical assays. The antibody transport probes 196 have a specific affinity for the target antigen 204. The transport probes 196 bind specifically to epitopes within the target antigen 204 as also shown in Fig. 8B. In one embodiment, antibody transport probes 196 which have an affinity for a portion of the target antigen may be conjugated to 3 micron magnetic capture beads 190 via EDC conjugation. Alternatively, conjugation of the transport probes 196 to the capture bead 190 may be achieved by passive adsorption.

With reference now to Step II shown in Fig. 11B, the target antigen 204 is added to the solution. The target antigen 204 binds to the antibody transport probe 196 attached to the capture bead 190 as also shown in Fig. 8B.

As illustrated in Step III, reporter beads 192 coated with antibody signal probes 208 are added to the solution. Antibody signal probes 208 specifically binds to the epitopes on target antigen 204 as also represented in Figs. 9B and 10B. In one embodiment, signal probes 208 are conjugated to 2.1 micron fluorescent reporter beads 192. Signal probes 208 and transport probes 196 each bind to specific epitopes on the target antigen, but not to each other. After adding reporter beads 192, the dual bead complex 194 is formed such that the target antigen 204 links capture bead 190 and reporter bead 192. With specific and thorough washing, there should be minimal non-specific binding between reporter bead 192 and capture bead 190.

In Step IV, after the binding in Step III, the dual bead complex 194 is separated from unbound reporter beads in the solution. The solution can be exposed to a magnetic field to capture the dual bead complex structures 194 using the magnetic properties of capture bead 190. The magnetic field can be encapsulated in a magnetic test tube rack 216 with a built-in magnet 218, which can be permanent or electromagnetic to draw out the magnetic beads and remove any

unbound reporter beads in the suspension. Note that capture beads not bound to reporter beads will also be isolated.

The purification process of Step IV includes the removal of supernatant containing free-floating particles. Wash buffer is added into the test tube and the
5 bead solution is mixed well. Most of the unbound reporter beads 182, free-floating protein samples, and non-specifically bound particles are agitated and removed from the supernatant. The dual bead complex can form a matrix of capture beads, target antigen, and reporter beads, wherein the wash process can further assist in the extraction of free floating particles trapped in the lattice structure of overlapping
10 dual bead particles.

The last principal step in Fig. 11B is Step V. In this step, once the dual bead complex has been washed approximately 3-5 times with wash buffer solution, the assay mixture is loaded into the disc and is thereby in condition to be analyzed.

Fig. 12A shows an alternative genetic assay method referred to here as a
15 "two-step hybridization" to create the dual bead complex which has 6 principal steps. Generally, capture beads are coated with oligonucleotide transport probes 198 complementary to DNA or RNA target agent and placed into a buffer solution. In this embodiment, transport probes which are complementary to a portion of target agent are conjugated to 3 micron magnetic capture beads via EDC conjugation.
20 Other type of conjugation of the oligonucleotide transport probes to a solid phase may be utilized. These include, for example, passive adsorption or use of streptavidin-biotin interactions. These 6 main steps according to this method of the present invention are consecutively identified as Steps I, II, III, IV, V, and VI in Fig. 12A. In summary, the steps and subsidiary steps of this method include providing a
25 mixture of capture beads having transport probes covalently attached thereto, providing a mixture of reporter beads that have signal probes covalently bound thereto, blocking the mixture of capture beads with a bead blocking agent, blocking the mixture of reporter beads with the bead blocking agent, suspending the mixture of capture beads in a hybridization solution, adding to the mixture a target agent that
30 hybridizes with the transport probes, allowing the transport probes to hybridize with the target agent to thereby form a hybridized partial complex including at least one capture bead, separating within the mixture the hybridized partial complex from unbound target agents, adding to the mixture reporter beads including signal probes covalently attached thereto, allowing the signal probes to hybridize with the target

agent to thereby form a dual bead complex including at least one capture bead and one reporter bead, separating the dual bead complex from unbound reporter beads, removing from the mixture the unbound reporter beads, and loading the mixture including the dual bead complex into an optical bio-disc for analysis.

5 More specifically now with reference to Step I shown in Fig. 12A, capture beads 190, suspended in hybridization solution, are loaded from the pipette 214 into the test tube 212. The preferred hybridization solution is composed of 0.2M NaCl, 10mM MgCl₂, 1mM EDTA, 50mM Tris-HCl, pH 7.5, and 5X Denhart's mix to serve as the blocking agent. Alternatively, salmon sperm DNA may be used as the
10 blocking agent. A desirable hybridization temperature is 37 degrees Celsius.

In Step II, target DNA or RNA 202 is added to the solution and binds to the complementary sequences of transport probe 198 attached to capture bead 190. In one specific embodiment of this method, target agent 202 and the transport probe 198 are allowed to hybridize for 2 to 3 hours at 37 degrees Celsius. Sufficient
15 hybridization, however, may be achieved within 30 minutes at room temperature. At higher temperatures, hybridization may be achieved substantially instantaneously.

As next shown in Step III, target agents 202 bound to the capture beads are separated from unbound species in solution by exposing the solution to a magnetic field to isolate bound target sequences by using the magnetic properties of the
20 capture bead 190. The magnetic field can be enclosed in a magnetic test tube rack 216 with a built-in magnet permanent 218 or electromagnet to draw out the magnetic beads and remove any unbound target DNA 202 free-floating in the suspension via pipette extraction of the solution. A wash buffer is added and the separation process can be repeated. The preferred wash buffer after the transport
25 probes 198 and target DNA 202 hybridize, consists of 145mM NaCl, 50mM Tris, pH 7.5, and 0.05% Tween. Hybridization methods and techniques for decreasing non-specific binding of target agents to beads are further disclosed in commonly assigned and co-pending U.S. Provisional Application Serial No. 60/278,691 entitled
"Reduction of Non-Specific Binding of Dual Bead Assays by Use of Blocking
30 Agents" filed March 26, 2001. This application is herein incorporated by reference in its entirety.

Referring now to Step IV illustrated in Fig. 12A, reporter beads 192 are added to the solution as discussed in conjunction with the method shown in Fig. 11A. As indicated above, the reporter beads are suspended in the hybridization buffer

including salmon sperm DNA or Denhart's solution. Reporter beads 192 are coated with signal probes 206 that are complementary to target agent 202. In one particular embodiment of this method, signal probes 206, which are complementary to a portion of target agent 202, are conjugated to 2.1 micron fluorescent reporter beads 192. Signal probes 206 and transport probes 198 each have sequences that are complementary to target agent 202, but not complementary to each other. After the addition of reporter beads 192, the dual bead complex structures 190 are formed. As would be readily apparent to one of skill in the art, the dual bead complex structures are formed only if the target agent of interest is present. In this formation, target agent 202 links magnetic capture bead 190 and reporter bead 192. Using the preferred buffer solution, with specific and thorough washing, there is minimal non-specific binding between the reporter beads and the capture beads. Target agent 202 and signal probe 206 are preferably allowed to hybridize for 2-3 hours at 37 degrees Celsius. As with Step II discussed above, sufficient hybridization may be achieved within 30 minutes at room temperature. At higher temperatures, the hybridization taking place in this step may also be achieved substantially instantaneously.

With reference now to Step V shown in Fig. 12A, after the hybridization in Step IV, the dual bead complex 194 is separated from unbound species in solution. The solution is again exposed to a magnetic field to isolate the dual bead complex 194 using the magnetic properties of the capture bead 190. Note again that the isolate will include capture beads not bound to reporter beads.

A purification process to remove supernatant containing free-floating particles includes adding wash buffer into the test tube and mixing the bead solution well. The preferred wash buffer for the two-step assay consists of 145mM NaCl, 50mM Tris, pH 7.5, 0.1% SDS, 0.05% Tween, 0.25% NFDM, and 10mM EDTA. Most unbound reporter beads, free-floating DNA, and non-specifically bound particles are agitated and removed from the supernatant. The dual bead complex can form a matrix of capture beads, target agents, and reporter beads, wherein the wash process can further assist in the extraction of free floating particles trapped in the lattice structure of overlapping dual bead particles. Other related aspects directed to reduction of non-specific binding between reporter bead, target agent, and capture bead are disclosed in, for example, commonly assigned and co-pending U.S. Provisional Application Serial No. 60/272,243 entitled "Mixing Methods to

Reduce Non-Specific Binding in Dual Bead Assays" filed February 28, 2001; and U.S. Provisional Application Serial No. 60/272,485 entitled "Dual Bead Assays Including Linkers to Reduce Non-Specific Binding" filed March 1, 2001, which are incorporated herein in their entirety.

5 The final principal step shown in Fig. 12A is Step VI. In this step, once the dual bead complex 194 has been washed approximately 3-5 times with wash buffer solution, the assay mixture is loaded into the disc and analyzed. Alternatively, during this step, the oligonucleotide signal and transport probes may be ligated to prevent breakdown of the dual bead complex during the disc analysis and signal
10 detection processes. Further details regarding probe ligation methods are disclosed in commonly assigned and co-pending U.S. Provisional Application Serial No. 60/278,694 entitled "Improved Dual Bead Assays Using Ligation" filed March 26, 2001, which is herein incorporated in its entirety by reference.

15 In accordance with another aspect of this invention, Fig. 12B shows an immunoassay method, similar to those discussed in connection with Figs. 11B and 12A, referred to here as a "two-step binding" to create the dual bead complex in an immunochemical assay. As with the method shown in Fig. 12A, this method includes 6 main steps. In general, capture beads coated with antibody transport probes which specifically binds to epitopes on target antigen are placed into a buffer
20 solution. In one specific embodiment, antibody transport probes are conjugated to 3 micron magnetic capture beads. Different sized magnetic capture beads may be employed depending on the type of disc drive and disc assembly utilized to perform the assay. These 6 main steps according to this alternative method of the invention are respectively identified as Steps I, II, III, IV, V, and VI in Fig. 12B.

25 With specific reference now to Step I shown in Fig. 12B, capture beads 190, suspended in buffer 210 solution, are loaded into a test tube 212 via injection from pipette 214. In this embodiment, the buffer solution 210 includes a blocking agent such as salmon sperm DNA, Denhart's solution, and BSA. The preferred blocking agent is BSA.

30 In Step II, target antigen 204 is added to the solution and binds to the antibody transport probe 196 attached to capture bead 190. Target antigen 204 and the transport probe 196 are preferably allowed to bind for 2 to 3 hours at 37 degrees Celsius. Shorter binding times are also possible.

As shown in Step III, target antigen 204 bound to the capture beads 190 are separated from unbound species in solution by exposing the solution to a magnetic field to isolate bound target proteins or glycoproteins by using the magnetic properties of the capture bead 190. The magnetic field can be enclosed in a magnetic test tube rack 216 with a built-in magnet permanent 218 or electromagnet to draw out the magnetic beads and remove any unbound target antigen 204 free-floating in the suspension via pipette extraction of the solution. A wash buffer is added and the separation process can be repeated.

As next illustrated in Step IV, reporter beads 192 are added to the solution as discussed in conjunction with the method shown in Fig. 11B. Reporter beads 192 are coated with signal probes 208 that have an affinity for the target antigen 204. In one particular embodiment of this two-step immunochemical assay, signal probes 208, which bind specifically to a portion of target agent 204, are conjugated to 2.1 micron fluorescent reporter beads 192. Signal probes 208 and transport probes 196 each bind to specific epitopes on the target agent 204, but do not bind to each other. After the addition of reporter beads 192, the dual bead complex structures 190 are formed. As would be readily apparent to those skilled in the art, these dual bead complex structures are formed only if the target antigen of interest is present. In this formation, target antigen 204 links magnetic capture bead 190 and reporter bead 192. Using the preferred buffer solution, with specific and thorough washing, there is minimal non-specific binding between the reporter beads and the capture beads. As with Step I above, the buffer solution 210 includes a blocking agent such as salmon sperm DNA, Denhart's solution, and BSA. The preferred blocking agent is BSA. Target antigen 204 and signal probe 208 are allowed to hybridize for 2-3 hours at 37 degrees Celsius. As with Step II discussed above, sufficient binding may be achieved within 30 minutes at room temperature. In the case of immunoassays temperatures higher than 37 degrees Celsius are not preferred because the proteins will denature.

Turning next to Step V as illustrated in Fig. 12B, after the binding shown in Step IV, the dual bead complex 194 is separated from unbound species in solution. This is achieved by exposing the solution to a magnetic field to isolate the dual bead complex 194 using the magnetic properties of the capture bead 190 as shown. Note again that the isolate will include capture beads not bound to reporter beads.

A purification process to remove supernatant containing free-floating particles includes adding wash buffer into the test tube and mixing the bead solution well. Most unbound reporter beads, free-floating proteins, and non-specifically bound particles are agitated and removed from the supernatant. The dual bead complex
 5 can form a matrix of capture beads, target agents, and reporter beads, wherein the wash process can further assist in the extraction of free floating particles trapped in the lattice structure of overlapping dual bead particles.

The final main step shown in Fig. 12B is Step VI. In this step, once the dual bead complex 194 has been washed approximately 3-5 times with wash buffer
 10 solution, the assay mixture is loaded into the disc and analyzed.

With reference now to Fig. 13, there is shown a cross sectional view illustrating the disk layers (similar to Fig. 6) of the mixing or loading chamber 164. Access to the loading chamber 164 is achieved by an inlet port 152 where the dual bead assay preparation is loaded into the disc system.

Fig. 14 is a view similar to Fig. 13 showing the mixing or loading chamber 164 with the pipette 214 injection of the dual bead complex 194 onto the disc. In this example, the complex includes reporters 192 and capture bead 190 linked together by the target DNA or RNA 202. The signal DNA 206 is illustrated as single stranded DNA complementary to the capture agent. The discs illustrated in Figs. 13
 15 and 14 may be readily adapted to other assays including the immunoassays and general molecular assays discussed above which employ, alternatively, proteins such as antigens or antibodies implemented as the transport probes, target agents, and signal probes accordingly.

Fig. 15A shows the flow channel 160 and the target or capture zone 170 after
 25 anchoring of dual bead complex 194 to a capture agent 220. The capture agent 220 in this embodiment is attached to the active layer 176 by applying a small volume of capture agent solution to the active layer 176 to form clusters of capture agents within the area of the target zone 170. In this embodiment, the capture agent includes biotin or BSA-biotin. Fig. 15A also shows reporters 192 and capture beads
 30 190 as components of a dual bead complex 194 as employed in the present invention. In this embodiment, anchor agents 222 are attached to the reporter beads 192. The anchor agent 222, in this embodiment, may include Streptavidin or Neutravidin. So when the reporter beads 192 come in close proximity to the capture agents 220, binding occurs between the anchor probe 222/206 and the capture

agent 220, via biotin-streptavidin interactions, thereby retaining the dual bead complex 194 within the target zone 170. At this point, an interrogation beam 224 directed to the target zone 170 can be used to detect the dual bead complex 194 within the target zone 170.

5 Fig. 15B is a cross sectional view similar to Fig. 15A illustrating the entrapment of the reporter bead 192 within the target zone 170 after a subsequent change in disc rotational speed. The change in rotational speed removes the capture beads 190 from the dual bead complex 194, ultimately isolating the reporter bead 192 in the target zone 170 to be detected by the interrogation or read beam
10 224.

Fig. 16A is a cross sectional view, similar to Fig. 15A, that illustrates an alternative embodiment to Fig. 15A wherein the signal probes 206 or an anchor agent 222, on the reporter beads 192, directly hybridizes to the capture agent 220. Fig. 16A shows the flow channel 160 and the target or capture zone 170 after
15 anchoring of dual bead complex 194 with the capture agent 220. The capture agent 220 in this embodiment is attached to the active layer 176 by applying a small volume of capture agent solution to the active layer 176 to form clusters of capture agents within the area of the target zone 170. Alternatively, the capture agent 220 may be attached to the active layer using an amino group that covalently binds to
20 the active layer 176. In this embodiment, the capture agent includes DNA. Fig. 16A also shows reporters 192 and capture beads 190 as components of a dual bead complex 194 as employed in the present invention. In this embodiment, anchor probes 222 are attached to the reporter beads 192. The anchor agent 222, in this embodiment, may be a specific sequence of nucleic acids that are complimentary to
25 the capture agent 220 or the oligonucleotide signal probe 206 itself. So when the reporter beads 192 come in close proximity to the capture agents 220, hybridization occurs between the anchor probe 222 and the capture agent 220 thereby retaining the dual bead complex 194 within the target zone 170. In an alternate embodiment, the signal probe 206 serves the function of anchor probe 222. At this point, an
30 interrogation beam 224 directed to the target zone 170 may be used to detect the dual bead complex 194 within the target zone 170.

Fig. 16B illustrates the embodiment in Fig. 16A after a subsequent change in disc rotational speed. The change in rational speed removes the capture bead 190 from the dual bead complex 194, ultimately isolating the reporter bead 192 and the

target DNA sequence 202 in the target zone 170 to be detected by an interrogation beam 224.

Referring now to Fig. 17, there is shown an alternative to the embodiment illustrated in Fig. 15A. In this embodiment, anchor agents 222 are attached to the capture beads 190 instead of the reporter beads. The anchor agent 222, in this embodiment, may include Streptavidin or Neutravidin. As in Fig. 15A, the target zone 170 is coated with a capture agent 220. The capture agent may include biotin or BSA-biotin. Fig. 17 also shows reporters 192 and capture beads 190 as components of a dual bead complex 194 as employed in the present invention. When the capture beads 190 come in close proximity to the capture agents 220, binding occurs between the anchor probe 222 and the capture agent 220, via biotin-streptavidin interactions, thereby retaining the dual bead complex 194 within the target zone 170. At this point, an interrogation beam 224 directed to the target zone 170 can be used to detect the dual bead complex 194 within the target zone 170.

Fig. 18 is an alternative to the embodiment illustrated in Fig. 16A. In this embodiment, anchor agents 222 are attached to the capture beads 190 instead of the reporter beads. In this embodiment the transport probes 198, or an anchor agent 222 on the capture bead 190, directly hybridizes to the capture agent 220. In this embodiment, the capture agent 220 includes specific sequences of nucleic acid. The anchor agent 222, in this embodiment, may be a specific sequence of nucleic acids that are complimentary to the capture agent 220 or the oligonucleotide signal transport probe 198 itself. So when the capture beads 190 come in close proximity to the capture agents 220, hybridization occurs between the anchor agent 222 and the capture agent 220 thereby retaining the dual bead complex 194 within the target zone 170. At this point, an interrogation beam 224 directed to the target zone 170 can be used to detect the dual bead complex 194 within the target zone 170.

Figs. 19A-19C are detailed partial cross sectional views showing the active layer 176 and the substrate 174 of the present bio-disc 110 as implemented in conjunction with the genetic assays discussed herein. Figs. 19A-19C illustrates the capture agent 220 attached to the active layer 176 by applying a small volume of capture agent solution to the active layer 176 to form clusters of capture agents within the area of the target zone. The bond between capture agent 220 and the active layer 176 is sufficient so that the capture agent 220 remains attached to the active layer 176 within the target zone when the disc is rotated. Figs. 19A and 19B

also depict the capture bead 190 from the dual bead complex 194 binding to the capture agent 220 in the capture zone. These dual bead complexes are prepared according to the methods such as those discussed in Figs. 11A and 12A. The capture agent 220 includes biotin and BSA-biotin. In this embodiment, the reporter bead 192 anchors the dual bead complex 194 in the target zone via biotin/streptavidin interactions. Alternatively, the target zone may be coated with streptavidin and may bind biotinylated reporter beads. Fig. 19C illustrates an alternative embodiment which includes an additional step to those discussed in connection with Figs. 19A and 19B. In this preferred embodiment, a variance in the disc rotations per minute may create a centrifugal force enough to break the capture beads 190 away from the dual bead complex 194 based on the differential size and/or mass of the bead. Although there is a shift in the rotation speed of the disc, the reporter bead 192 remains anchored to the target zone. Thus, the reporter beads 192 are maintained within the target zone and detected using an optical bio-disc reader.

Figs. 20A, 20B, and 20C illustrate an alternative embodiment to the embodiment discussed in Figs. 19A-19C. Figs. 20A-20C show detailed partial cross sectional views of a target zone implemented in conjunction with immunochemical assays. Figs. 20A and 20B also depict the capture bead 190 from the dual bead complex 194 binding to the capture agent 220 in the capture zone. The capture agent 220 includes biotin and BSA-biotin. These dual bead complexes may be prepared according to methods such as those discussed in Figs. 11B and 12B. In this embodiment, the reporter bead 192 anchors the dual bead complex 194 in the target zone via biotin/streptavidin interactions.

Referring now to Figs. 21A, 21B, and 21C, there is shown detailed partial cross sectional views of a target zone including the active layer 176 and the substrate 174 of the present bio-disc 110 as implemented in conjunction with the genetic assays discussed herein. Figs. 21A-21C illustrate the capture agent 220 attached to the active layer 176 by use of an amino group 226 which is made an integral part of the capture agent 220. As indicated, the capture agent 220 is situated within the target zone. The bond between the amino group 226 and the capture agent 220, and the amino group 226 and the active layer 176 is sufficient so that the capture agent 220 remains attached to the active layer 176 within the target zone when the disc is rotated. The preferred amino group 226 is NH_2 . A thiol group

may alternatively be employed in place of the amino group 226. In this embodiment of the present invention, the capture agent 220 includes the specific sequences of amino acids that are complimentary to the anchor agent 222 or oligonucleotide signal probe 206 which are attached to the reporter bead 192.

5 Fig. 21B depicts the reporter bead 192 of the dual bead complex 194, prepared according to methods such as those discussed in Figs. 11A and 12A, binding to the capture agent 220 in the target zone. As the dual bead complex 194 flows towards the capture agent 220 and is in sufficient proximity thereto, hybridization occurs between the anchor agent 222 or oligonucleotide signal probe
10 206 and the capture agent 220. Thus, the reporter bead 192 anchors the dual bead complex 192 within the target zone.

 Fig. 21C illustrates an alternative embodiment that includes an additional step to those discussed in connection with Figs. 21A-21B. In this preferred embodiment, a variance in the disc rotations per minute may create enough
15 centrifugal force to break the capture beads 190 away from the dual bead complex 194 based on the differential size and/or mass of the bead. Although there is a shift in the rotation speed of the disc, the reporter bead 192 with the target DNA sequence 202 remains anchored to the target zone. In either case, the reporter beads 192 are maintained within the target zone as desired.

20 Figs. 22A, 22B, and 22C illustrate an alternative embodiment to the embodiment discussed in Figs. 21A-21C. Figs. 22A-22C show detailed partial cross sectional views of a target zone implemented in conjunction with immunochemical assays. Figs. 22A and 22B also depict the reporter bead 192 from the dual bead complex 194, prepared according to methods such as those discussed in Figs. 11B
25 and 12B, binding to the capture agent 220 in the capture zone. In this embodiment, the capture agent 220 includes antibodies bound to the target zone by use of an amino group 226 which is made an integral part of the capture agent 220. Alternatively, the capture agents 220 may be bound to the active layer 176 by passive absorption, and hydrophobic or ionic interactions. In this embodiment, the
30 reporter bead 192 anchors the dual bead complex 194 in the target zone via specific antibody binding. As with the embodiment illustrated in Fig. 21C, Fig. 22C shows an alternative embodiment that includes an additional step to those discussed in connection with Figs. 22A-22B. In this alternative embodiment, a variance in the disc rotations per minute may create enough centrifugal force to break the capture

beads 190 away from the dual bead complex 194 based on the differential size and/or mass of the bead. Although there is a shift in the rotation speed of the disc, the reporter bead 192 with the target antigen 204 remains anchored to the target zone. In either case, the reporter beads 192 are maintained within the target zone as desired.

Figs. 23A and 23B are detailed partial cross sectional views showing the active layer 176 and the substrate 174 of the present bio-disc 110 as implemented in conjunction with the genetic assays. Figs. 23A and 23B illustrate an alternative embodiment to that discussed in Figs. 19A and 19B above. In contrast to the embodiment in Figs. 19A and 19B, in the present embodiment, the anchor agent 222 is attached to the capture bead 190 instead of the reporter bead 192. Fig. 23B illustrates the capture bead 190, from the dual bead complex 194, binding to the capture agent 220 in the capture zone. The capture agent 220 includes biotin and BSA-biotin. In this embodiment, the capture bead 190 anchors the dual bead complex 194 in the target zone via biotin/streptavidin interactions.

With reference now to Figs. 24A and 24B, there is presented detailed partial cross sectional views showing the active layer 176 and the substrate 174 of the present bio-disc 110 as implemented in conjunction with the genetic assays. Figs. 23A and 23B illustrate an alternative embodiment to that discussed in Figs. 21A and 21B above. In contrast to the embodiment in Figs. 21A and 21B, in the present embodiment, the anchor agent 222 is attached to the capture bead 190 instead of the reporter bead 192. Fig. 23B illustrates the capture bead 190, from the dual bead complex 194, binding to the capture agent 220 in the capture zone. The capture agent 220 is attached to the active layer 176 by use of an amino group 226 which is made an integral part of the capture agent 220. As indicated, the capture agent 220 is situated within the target zone. The bond between the amino group 226 and the capture agent 220, and the amino group 226 and the active layer 176 is sufficient so that the capture agent 220 remains attached to the active layer 176 within the target zone when the disc is rotated. In this embodiment of the present invention, the capture agent 220 includes the specific sequences of amino acids that are complimentary to the anchor agent 222 or oligonucleotide transport probe 198 which are attached to the capture bead 190. In this embodiment, the capture bead 190 anchors the dual bead complex 194 in the target zone via hybridization between the capture agent 220 and the anchor agent or the transport probe 198.

Disc Processing Methods

Turning now to Figs. 25A–25D, there is shown the target zones 170 set out in Figs. 21A-21C and Figs. 24A-24B in the context of a disc, using as an input the solution created according to methods such as those shown in Figs. 11A and 12A.

Fig. 25A shows a mixing/loading chamber 164, accessible through an inlet port 152, and leading to a flow channel 160. Flow channel 160 is pre-loaded with capture agents 220 situated in clusters in target zones 170. Each of the clusters of capture agents 220 is situated within a respective target zone 170. Each target zone 170 can have one type of capture agent or multiple types of capture agents, and separate target zones can have one and the same type of capture agent or multiple different capture agents in multiple capture fields. In the present embodiment, the capture agent can include specific sequences of nucleic acids that are complimentary to anchor agents 222 on either the reporter 192 or capture bead 190.

In Fig. 25B, a pipette 214 is loaded with a test sample of DNA or RNA that has been sequestered in the dual bead complex 194. The dual bead complex is injected into the flow channel 160 through inlet port 152. As flow channel 160 is further filled with the dual bead complex from pipette 214, the dual bead complex 194 begins to move down flow channel 160 as the disc is rotated. The loading chamber 164 can include a break-away retaining wall 228 so that complex 194 moves down the flow channel at one time.

In this embodiment, anchor agents 222, attached to reporter beads 192, bind to the capture agents 220 by hybridization, as illustrated in Fig. 25C. In this manner, reporter beads 192 are retained within target zone 170. Binding can be further facilitated by rotating the disc so that the dual bead complex 194 can slowly move or tumble down the flow channel. Slow movement allows ample time for additional hybridization. After hybridization, the disc can be rotated further at the same speed or faster to clear target zone 170 of any unattached dual bead complex 194, as illustrated in Fig. 25D.

An interrogation beam 224 can then be directed through target zones 170 to determine the presence of reporters, capture beads, and dual bead complex, as illustrated in Fig. 25D. In the event no target DNA or RNA is present in the test sample, there will be no dual bead complex structures, reporters, or capture beads

bound to the target zones 170, but a small amount of background signal may be detected in the target zones from non-specific binding. In this case, when the interrogation beam 224 is directed into the target zone 170, a zero or low reading results, thereby indicating that no target DNA or RNA was present in the sample.

5 The speed, direction, and stages of rotation, such as one speed for one period followed by another speed for another period, can all be encoded in the operational information on the disc.

10 Figs. 26A–26D show the target zones 170 including the capture chemistries discussed in Figs. 19A–19C and Figs. 23A–23B. This method uses as an input, the solution created according to methods shown in Figs. 11A and 12A. Figs. 26A–26D illustrate an alternative embodiment to that discussed in Figs. 25A–25D showing a different bead capture method described in further detail below.

15 Fig. 26A shows a mixing/loading chamber 164, accessible through an inlet port 152, and leading to a flow channel 160. Flow channel 160 is pre-loaded with capture agents 220 situated in clusters in target zones 170. Each of the clusters of capture agents 220 is situated within a respective target zone 170. Each target zone 170 can have one type of capture agent or multiple types of capture agents, and separate target zones can have one and the same type of capture agent or multiple different capture agents in multiple capture fields. In the present embodiment, the capture agent can include specific biotin and BSA-biotin that has affinity to the anchor agents 222 on either the reporter 192 or capture bead 190. The anchor agents may include Streptavidin and Neutravidin.

20 In Fig. 26B, a pipette 214 is loaded with a test sample of DNA or RNA that has been sequestered in the dual bead complex 194. The dual bead complex is injected into the flow channel 160 through inlet port 152. As flow channel 160 is further filled with the dual bead complex from pipette 214, the dual bead complex 194 begins to move down flow channel 160 as the disc is rotated. The loading chamber 164 can include a break-away retaining wall 228 so that complex 194 moves down the flow channel at one time.

30 In this embodiment, anchor agents 222, attached to reporter beads 192, bind to the capture agents 220 by biotin-streptavidin interactions, as illustrated in Fig. 26C. In this manner, reporter beads 192 are retained within target zone 170. Binding can be further facilitated by rotating the disc so that the dual bead complex 194 can slowly move or tumble down the flow channel. Slow movement allows

ample time for additional binding between the capture agent 220 and the anchor agent 222. After binding, the disc can be rotated further at the same speed or faster to clear target zone 170 of any unattached dual bead complex 194, as illustrated in Fig. 26D.

5 An interrogation beam 224 can then be directed through target zones 170 to determine the presence of reporters, capture beads, and dual bead complex, as illustrated in Fig. 26D. In the event no target DNA is present in the test sample, there will be no dual bead complex structures beads bound to the target zones 170. A small amount of background signal may be detected in the target zones from non-
10 specific binding. In this case, when the interrogation beam 224 is directed into the target zone 170, a zero or low reading results, thereby indicating that no target DNA or RNA was present in the sample.

 The speed, direction, and stages of rotation, such as one speed for one period followed by another speed for another period, can all be encoded in the
15 operational information on the disc.

 Referring next to Figs. 27A-27D the is shown a series of cross sectional side views illustrating the steps of yet another alternative method according to the present invention. Figs. 27A-27D show the target zones 170 including the capture mechanisms discussed in connection with Figs. 22A-22C. This method uses an
20 input the solution created according to the preparation methods shown in Figs. 11B and 12B. Figs. 27A-27D illustrate an immunochemical assay and an alternative bead capture method.

 Fig. 27A shows a mixing/loading chamber 164, accessible through an inlet port 152, and leading to a flow channel 160. Flow channel 160 is pre-loaded with
25 capture agents 220 situated in clusters in target zones 170. Each of the clusters of capture agents 220 is situated within a respective target zone 170. Each target zone 170 can have one type of capture agent or multiple types of capture agents, and separate target zones can have one and the same type of capture agent or multiple different capture agents in multiple capture fields. In the present
30 embodiment, the capture agent can include antibodies that specifically bind to epitopes on the anchor agents 222 on either the reporter 192 or capture bead 190. Alternatively, the capture agent can directly bind to epitopes on the target antigen 204 within the dual bead complex 194. The anchor agents can include the target antigen, antibody transport probe 196, the antibody signal probe 208, or any

antigen, bound to either the reporter bead 192 or the capture bead 190, that has epitopes than can specifically bind to the capture agent 220.

In Fig. 27B, a pipette 214 is loaded with a test sample of target antigen that has been sequestered in the dual bead complex 194. The dual bead complex is injected into the flow channel 160 through inlet port 152. As flow channel 160 is further filled with the dual bead complex from pipette 214, the dual bead complex 194 begins to move down flow channel 160 as the disc is rotated. The loading chamber 164 may include a break-away retaining wall 228 so that complex 194 moves down the flow channel at one time.

In this embodiment, anchor agents 222, attached to reporter beads 192, bind to the capture agents 220 by antibody-antigen interactions, as illustrated in Fig. 27C. In this manner, reporter beads 192 are retained within target zone 170. Binding can be further facilitated by rotating the disc so that the dual bead complex 194 can slowly move or tumble down the flow channel. Slow movement allows ample time for additional binding between the capture agents 220 and the anchor agent 222. After binding, the disc can be rotated further at the same speed or faster to clear target zone 170 of any unattached dual bead complex 194, as illustrated in Fig. 27D.

An interrogation beam 224 can then be directed through target zones 170 to determine the presence of reporters, capture beads, and dual bead complex, as illustrated in Fig. 27D. In the event no target antigen is present in the test sample, there will be no dual bead complex structures, reporters, or capture beads bound to the target zones 170, but a small amount of background signal may be detected in the target zones from non-specific binding. In this case, when the interrogation beam 224 is directed into the target zone 170, a zero or low reading results, thereby indicating that no target was present in the sample.

The speed, direction, and stages of rotation, such as one speed for one period followed by another speed for another period, can all be encoded in the operational information on the disc.

The methods described in Figs. 25A-25D, 26A-26D, and 27A-27D are implemented using the reflective disc system 144. It should be understood that these methods and any other bead or sphere detection may also be carried out using the transmissive disc embodiment 180, as described in Figs. 4A-4C, 5B, and 6B. It should also be understood that the methods described in Figs. 11A-11B, 12A-

12B, 25A-25D, 26A-26D, and 27A-27D are not limited to creating the dual bead complexes outside of the optical bio-discs but may include embodiments that use "in-disc" or "on-disc" formation of the dual bead complexes. In these on-disc implementations the dual bead complex is formed within the fluidic circuits of the optical bio-disc 110. For example, the dual bead formation may be carried out in the loading or mixing chamber 164. In one embodiment, the beads and sample are added to the disc at the same time, or nearly the same time. Alternatively, the beads with the probes can be pre-loaded on the disc for future use with a sample so that only a sample needs to be added.

10 The beads would typically have a long shelf life, with less shelf life for the probes. The probes can be dried or lyophilized (freeze dried) to extend the period during which the probes can remain in the disc. With the probes dried, the sample essentially reconstitutes the probes and then mixes with the beads to produce dual bead complex structures can be performed.

15 In either case, the basic process for on disc processing includes: (1) inserting the sample into a disc with beads with probes; (2) causing the sample and the beads to mix on the disc; (3) isolating, such as by applying a magnetic field, to hold the dual bead complex and move the non-held beads away, such as to a region referred to here as a waste chamber; and (4) directing the dual bead complexes (and any other material not moved to the waste chamber) to the capture fields. The detection process can be the same as one of those described above, such as by event detection or fluorimetry.

Detection and Related Signal Processing Methods and Apparatus

25 The number of reporter beads bound in the capture field can be detected in a qualitative manner, and may also be quantified by the optical disc reader.

The test results of any of the test methods described above can be readily displayed on monitor 114 (Fig. 1). The disc according to the present invention preferably includes encoded software that is read to control the controller, the processor, and the analyzer as shown in Fig. 2. This interactive software is implemented to facilitate the methods described herein and the display of results.

Fig. 28A is a graphical representation of an individual 2.1 micron reporter bead 192 and a 3 micron capture bead 190 positioned relative to the tracks A-E of an optical bio-disc according to the present invention.

Fig. 28B is a series of signature traces, from tracks A-E, derived from the beads of Fig. 28A utilizing a detected signal from the optical drive according to the present invention. These graphs represent the detected return beam 124. As shown, the signatures for a 2.1 micron reporter bead 190 are sufficiently different from those for a 3 micron capture bead 192 such that the two different types of beads can be detected and discriminated. A sufficient deflection of the trace signal from the detected return beam as it passes through a bead is referred to as an event.

Fig. 29A is a graphical representation of a 2.1 micron reporter bead and a 3 micron capture bead linked together in a dual bead complex positioned relative to the tracks A-E of an optical bio-disc according to the present invention.

Fig. 29B is a series of signature traces, from tracks A-E, derived from the beads of Fig. 29A utilizing a detected signal from the optical drive according to the present invention. These graphs represent the detected return beam 124. As shown, the signatures for a 2.1 micron reporter bead 190 are sufficiently different from those for a 3 micron capture bead 192 such that the two different types of beads can be detected and discriminated. A sufficient deflection of the trace signal from the detected return beam as it passes through a bead is referred to as an event. The relative proximity of the events from the reporter and capture bead indicates the presence or absence the dual bead complex. As shown, the traces for the reporter and the capture bead are right next to each other indicating the beads are joined in a dual bead complex.

Alternatively, other detection methods can be used. For example, reporter beads can be fluorescent or phosphorescent. Detection of these reporters can be carried out in fluorescent or phosphorescent type optical disc readers. Other signal detection methods are described, for example, in commonly assigned co-pending U.S. Patent Application Serial No. 10/008,156 entitled "Disc Drive System and Methods for Use with Bio-Discs" filed November 9, 2001, which is expressly incorporated by reference; U.S. Provisional Application Serial Nos. 60/270,095 filed February 20, 2001 and 60/292,108, filed May 18, 2001; and the above referenced U.S. Patent Application Serial No. 10/043,688 entitled "Optical Disc Analysis System Including Related Methods For Biological and Medical Imaging" filed January 10, 2002.

Fig. 30A is a bar graph of data generated using a fluorimeter showing a concentration dependent target detection using fluorescent reporter beads. This graph shows the molar concentration of target DNA versus number of detected beads. The dynamic range of target detection shown in the graph is $10E-16$ to $10E-10$ Molar (moles/liter). While the particular graph shown was generated using data from a fluorimeter, the results may also be generated using a fluorescent type optical disc drive.

Fig. 30B presents a standard curve demonstrating that the sensitivity of a fluorimeter is approximately 1000 beads in a fluorescent dual bead assay. The sensitivity of any assay depends on the sensitivity of the assay itself and on the sensitivity of the detection system. Referring to Figs. 30A-30C, various studies were done to examine the sensitivity of the dual bead assay using different detection methods, e.g., a fluorimeter, and bio-disc detection according to the present invention.

As shown in Fig. 30B, the sensitivity of a fluorimeter is approximately 1000 beads in a fluorescent dual bead assay. In contrast, Fig. 30A shows that even at $10E-16$ Molar (moles/liter), a sufficient number of beads over zero concentration can be detected to sense the presence of the target. With a sensitivity of $10E-16$ Molar, a dual bead assay represents a very sensitive detection method for DNA that does not require DNA amplification (such as through PCR) and can be used to detect even a single bead.

In contrast to conventional detection methods, the use of a bio-disc coupled with a CD-reader or optical bio-disc drive (Fig. 1) improves the sensitivity of detection. For example, while detection with a fluorimeter is limited to approximately 1000 beads (Fig. 30B), use of a bio-disc coupled with CD-reader may enable the user to detect a single bead with the interrogation beam (Fig. 30C). Thus, the bioassay system provided herein improves the sensitivity of dual bead assays significantly. The detection of single beads using an optical bio-disc is discussed in detail in conjunction with Figs. 28A and 28B above. Fig. 28B shows the signal traces of each bead as detected by the bio-disc reader. Dual bead complexes may also be identified by the bio-disc reader using the unique signature trace collected from the detection of a dual bead complex as shown in Figs. 29A and 29B. Different optical bio-disc platforms may be used in conjunction with the reader device for

detection of beads including a reflective and a transmissive disc format illustrated in Figs. 3C and 4C, respectively.

Fig. 30C is a pictorial demonstrating the formation of the dual bead complex linked together by the presence of the target in a genetic assay. Sensitivity to within
 5 one reporter molecule is possible with the present dual bead assay quantified with a bio-CD reader shown in Figs. 1 and 2 above. Similarly, the dual bead complex formation may also be implemented in an immunochemical assay format as illustrated in Figs. 7B, 8B, 9B, 10B, 11B, and 12B above.

Fig. 31 shows data generated using a fluorimeter illustrating the
 10 concentration dependent detection of two different targets. The target detection was carried out in two different methods, the single and the duplex assays. In the single assay, the capture bead contains a transport probe specific to a single target and a reporter probe coated with a signal probe specific to the same target is mixed in a solution together with the target. In the duplex assay, the capture bead contains
 15 two different transport probes specific to two different targets. Experimental details regarding the use of the duplex target detection method is discussed in further detail in Example 2. Mixing different reporter beads (red and green fluorescent or silica and polystyrene beads, for example) containing signal probes specific to one of the two targets, allows the detection of two different targets simultaneously.

20 Detection of the dual bead duplex assay may be carried out using a magneto optical disc system described below. Figs. 32 and 37 illustrate the formation and binding of various dual bead complexes onto an optical disc which may be detected by an optical bio-disc drive (Fig. 2), a magneto-optical disc system, a fluorescent disc system, or any similar device. Unique signature traces of a dual bead complex
 25 collected from an optical disc reader are shown in Fig. 29B above. The traces from Fig. 29B further illustrate that different bead types can be detected by an optical disc reader since different type beads will show a different signature profile.

Multiplexing, Magneto-Optical, and Magnetic Discs Systems

30 The use of a dual bead assay in the capture of targets allows for the use in multiplexing assays. This type of multiplexing is achieved by combining different sizes of magnetic beads and different types and sizes of reporter beads, different target agents can be detected simultaneously. As indicated in Fig. 32, four sizes of magnetic capture beads, and four sizes of three types of reporter beads produce up

to 48 different types of dual bead complex. In a multiplexing assay, probes specific to different targets are thus conjugated to capture beads and reporter beads having different physical and/or optical properties, such as fluorescence at different wavelengths, to allow for the detection of different target agents simultaneously from the same biological sample in the same assay. As indicated in Figs. 16A, 16B, 17A, and 17B, small differences in size can be detected by detecting reflected or transmitted light.

Multiple dual bead complex structures to capture different target agents can be carried out on or off the disc. If off the disc, the dual bead suspension is loaded into a port on the disc. The port is sealed and the disc is rotated in the disc reader. During spinning, free (unbound) beads are spun off to a periphery of the disc. The reporter beads detecting various target agents are thus localized in capture fields. In this manner, the presence of a specific target agent can be detected, and the amount of a specific target agent can be quantified by the disc reader.

Fig. 33A is a general representation of an optical disc according to this aspect of the present invention and a method corresponding generally to the single-step method of Fig. 11A and 11B is shown. The sample and beads can be added at one time or successively but closely in time, or the beads can be pre-loaded into a portion of the disc. These materials can be provided to a mixing chamber 164 that can have a breakaway wall 228 (see Fig. 25A) that holds in the solution within the mixing chamber 164. Mixing the sample and beads on the disc would be accomplished through rotation at a rate insufficient to cause the wall to break or the capillary forces to be overcome.

The disc can be rotated in one direction, or it can be rotated alternately in opposite directions to agitate the material in a mixing chamber. The mixing chamber is preferably sufficiently large so that circulation and mixing is possible. The mixing can be continuous or intermittent.

Fig. 33B shows one embodiment of a rotationally directionally dependent valve arrangement that is directionally dependent and uses a movable component for a valve. The mixing chamber leads to an intermediate chamber 244 that has a movable component, such as a ball 246. In the non-rotated state, the ball 246 may be kept in a slight recessed portion, or chamber 244 may have a gradual V-shaped tapering in the circumferential direction to keep the ball centered when there is no rotation.

Referring to Figs. 33C and 33D in addition to Figs. 33A and 33B, when the disc is rotated clockwise (Fig. 33C), ball 246 moves to a first valve seat 248 to block passage to detection chamber 234 and to allow flow to waste chamber 232, shown in Fig. 33A. When the disc is rotated counter-clockwise (FIG. 33D), ball 246 moves
 5 to a second valve seat 250 to block a passage to waste chamber 232 and to allow flow to detection chamber 234.

Figs. 34A-34C show a variation of the prior embodiment in which the ball is replaced by a wedge 252 that moves one way or the other in response to acceleration of the disc. The wedge 252 can have a circular outer shape that
 10 conforms to the shape of an intermediate chamber 244. The wedge is preferably made of a heavy dense material relative to chamber 244 to avoid sticking. A coating can be used to promote sliding of the wedge relative to the chamber.

When the disc is initially rotated clockwise (Fig. 34B), the angular acceleration causes wedge 252 to move to block a passage to detection chamber
 15 234 and to allow flow to waste chamber 232. When the disc is initially rotated counter-clockwise (Fig. 34C), the angular acceleration causes wedge 252 moves to block a passage to waste chamber 232 and allow flow to detection chamber 234. During constant rotation after the acceleration, wedge 252 remains in place blocking the appropriate passage.

20 In another embodiment of the present invention where the capture beads are magnetic, a magnetic field from a magnetic field generator or field coil 230 can be applied over the mixing chamber 164 to hold the dual bead complexes and unbound magnetic beads in place while material without magnetic beads are allowed to flow away to a waste chamber 232. This technique may also be employed to aid in
 25 mixing of the assay solution within the fluidic circuits or channels before any unwanted material is washed away. At this stage, only magnetic capture beads, unbound or as part of a dual bead complex, remain. The magnetic field is released, and the dual bead complex with the magnetic beads is directed to a capture and detection chamber 234.

30 The process of directing non-magnetic beads to waste chamber 232 and then magnetic beads to capture chamber 234 can be accomplished through the microfluidic construction and/or fluidic components. A flow control valve 236 or some other directing arrangement can be used to direct the sample and non-magnetic beads to waste chamber 232 and then to capture chamber 234. A

number of embodiments for rotationally dependent flow can be used. Further details relating to the use of flow control mechanisms are disclosed in commonly assigned co-pending U.S. Patent Application Serial No. 09/997,741 entitled "Dual Bead Assays Including Optical Biodiscs and Methods Relating Thereto" filed
 5 November 27, 2001, which is herein incorporated by reference in its entirety.

Fig. 35 is a perspective view of a disc including one embodiment of a fluidic circuit employed in conjunction with magnetic beads and the magnetic field generator 230 according to the present invention. Fig. 35 also shows the mixing chamber 164, the waste chamber 232, and the capture chamber 234. The magnetic
 10 field generator 230 is positioned over disc 110 and has a radius such that as disc 110 rotates, magnetic field generator 230 remains over mixing chamber 164, and is radially spaced from chambers 232 and 234. As with the prior embodiment discussed above, a magnetic field from the magnetic field generator 230 can be applied over the mixing chamber 164 to hold the dual bead complexes and/or
 15 unbound magnetic beads in place while additional material is allowed to enter the mixing chamber 164. The method of rotating the disc while holding magnetic beads in place with the magnetic field generator 230 may also be employed to aid in mixing of the assay solution within the mixing chamber 164 before the solution contained therein is directed elsewhere.

20 Figs. 36A-36C are plan views illustrating a method of separation and detection for dual bead assays using the fluidic circuit shown in Fig. 35. With specific reference to Fig. 36A, there is shown an unrotated optical disc with a mixing chamber 164 shaped as an annular sector holds a sample with dual bead complexes 194 and various unbound reporter beads 192. The electromagnet is
 25 activated and the disc is rotated counter-clockwise (Fig. 36B), or it can be agitated at a lower rpm, such as 1X or 3X. Dual bead complexes 194, with magnetic capture beads, remain in mixing chamber 164 while the liquid sample and the unbound reporter beads 192 move in response to angular acceleration to a rotationally trailing end of mixing chamber 164. The disc is rotated with sufficient speed to overcome
 30 capillary forces to allow the unbound reporter beads in the sample to move through a waste fluidic circuit 238 to waste chamber 232. At this stage in the process, the liquid will not move down the capture fluidic circuit 240 because of the physical configuration of the fluidic circuit as illustrated.

As illustrated next in Fig. 36C, the magnet is deactivated and the disc is rotated clockwise. Dual bead complexes 194 move to the opposite trailing end of the mixing chamber 164 in response to angular acceleration and then through a capture fluidic circuit 240 to the capture chamber 234. At this later stage in the process, the dual bead solution will not move down the waste fluidic circuit 238 due to the physical layout of the fluidic circuit, as shown. The embodiment shown in Figs. 36A-36C thus illustrates directionally dependent flow as well as rotational speed dependent flow.

In this embodiment and others in which a fluidic circuit is formed in a region of the disc, a plurality of regions can be formed and distributed about the disc, for example, in a regular manner to promote balance. Furthermore, as discussed above, instructions for controlling the rotation can be provided on the disc. Accordingly, by reading the disc, the disc drive can have instructions to rotate for a particular period of time at a particular speed, stop for some period of time, and rotate in the opposite direction for another period of time. In addition, the encoded information can include control instructions such as those relating to, for example, the power and wavelength of the light source. Controlling such system parameters is particularly relevant when fluorescence is used as a detection method.

In yet another embodiment, a passage can have a material or configuration that can seal or dissolve either under influence from a laser in the disc drive, or with a catalyst pre-loaded in the disc, or such a catalyst provided in the test sample. For example, a gel may solidify in the presence of a material over time, in which case the time to close can be set sufficiently long to allow the unbound capture beads to flow to a waste chamber before the passage to the waste chamber closes. Alternatively, the passage to the waste chamber can be open while the passage to the detection chamber is closed. After the unbound beads are directed to the waste chamber, the passage to the detection chamber is opened by energy introduced from the laser to allow flow to the detection chamber.

Fig. 37 illustrates yet another embodiment of the optical disc 110 for use with a multiplexing dual bead assay. In this case, a disc, such as one used with a magneto-optical drive, has magnetic regions 242 that can be written and erased with a magnetic head. A magneto-optical disc drive, for example, can create magnetic regions 242 as small as 1 micron by 1 micron square. The close-up

section of the magnetic region 242 shows the direction of the magnetic field with respect to adjacent regions.

The ability to write to small areas in a highly controllable manner to make them magnetic allows capture areas to be created in desired locations. These
 5 magnetic capture areas can be formed in any desired configuration or location in one chamber or in multiple chambers. These areas capture and hold magnetic beads when applied over the disc. The domains can be erased if desired, thereby allowing them to be made non-magnetic and allowing the beads to be released.

In one configuration of a magnetic bead array according to this aspect of the
 10 present invention, a set of three radially oriented magnetic capture regions 244 are shown, by way of example, with no beads attached to the magnetic capture regions in the columns. With continuing reference to Fig. 37, there is shown a set of four columns in Section A with individual magnetic beads magnetically attached to the magnetic areas in a magnetic capture region. Another set of four columns arrayed
 15 in Section B is shown after binding of reporter beads to form dual bead complexes attached to specific magnetic areas, with different columns having different types of reporter beads. As illustrated in Section B, some of the reporter beads utilized vary in size to thereby achieve the multiplexing aspects of the present invention as implemented on a magneto-optical biodisc. In Section C, a single column of various
 20 dual bead complexes is shown as another example of multiplexing assays employing various bead sizes individually attached at separate magnetic areas.

In a method for use with such a magneto-optical biodisc, the write head in an MO drive can be used to create magnetic areas, and then a sample can be directed over that area to capture magnetic beads provided in the sample. After introduction
 25 of the first sample set, other magnetic areas may also be created and another sample set can be provided to the newly created magnetic capture region for detection. Thus detection of multiple sample sets may be performed on a single disc at different time periods. The magneto-optical drive also allows the demagnetization of the magnetic capture regions to thereby release and isolate the magnetic beads if
 30 desired. Thus this system provides for the controllable capture, detection, isolation, and release of one or more specific target molecules from a variety of different biochemical, chemical, or biological samples.

As described above, a sample can be provided to a chamber on a disc. Alternatively, a sample could be provided to multiple chambers that have sets of

With such a disc, a large number of tests can be performed at one time and can be performed interactively. In this manner, when a test is performed and a result is obtained, the system can be instructed to create a new set of magnetic regions for capturing the dual bead complex. Regions can be created one at a time or in large groups, and can be performed in successive chambers that have different pre-loaded beads. Other processing advantages can be obtained with a disc that has writeable magnetic regions. For example, the "capture agent" is essentially the magnetic field created by in the magnetic region on the disc and therefore there is no need to add an additional biological or chemical capture agent.

Methods for Decreasing Non-Specific Bead Binding

Fig. 39 is a bar graph of data generated using a fluorimeter showing a concentration dependent target detection using fluorescent reporter beads in a dual

bead assay. This graph shows the picomolar concentration of target DNA versus number of beads bound in a dual bead complex. The dynamic range of target detection shown in this graph is 0.25 pM to 2500 pM (picomoles/liter). While this particular graph was generated using data from a fluorimeter, the results may also
 5 be generated using a fluorescent type optical disc drive. Experimental details from an experiment related to detection of a range of target concentrations is discussed in detail in Example 3.

Referring now to Figs. 40, 41, 42, 43A, and 43B, here are shown data from experiments performed to determine the optimal concentration or ionic strength of
 10 various salts in the hybridization buffer or assay buffer. The salt concentration in the assay buffer needs to be optimized in order to increase hybridization efficiency or binding efficiency and decrease non-specific bead binding between capture and reporter beads resulting in lower signal to noise ratio which increases the sensitivity of the assay. In general, the data presented in these figures show that 40mM
 15 EDTA, 300mM NaCl, 30mM MgCl² are the optimal salt concentrations for use in one embodiment of the dual bead assay.

With specific reference to Fig. 40, there is a bar graph presentation showing data collected from an experiment using various concentrations of NaCl in the bead buffer and the related non-specific binding as a result of changes in the ionic
 20 strength of the buffer. Based on the results presented in Fig. 40, the optimal bead buffer concentration of sodium chloride for use in the dual bead assay is 0.2M since non-specific bead binding is minimal at this NaCl concentration. A detailed description of the experimental procedure used to generate this data is discussed below in Example 6.

Now referring to Fig. 41, there is shown a bar graph illustrating the effect of increasing EDTA concentration on the dual bead assay sensitivity using different target concentrations. Fig. 41 also shows the related non-specific binding as
 25 affected by the concentration of EDTA in the assay buffer. The optimal EDTA buffer concentration, based on the data presented, for use in the hybridization buffer is 40mM since signal generated from the dual bead assay was highest at this
 30 concentration.

Similarly Fig. 42 presents a bar graph presentation showing the effect of increasing NaCl concentration on the dual bead assay sensitivity using different target concentrations. The non-specific bead binding data related to optimization of

the buffer concentration of NaCl is represented in Fig. 40. As shown in Fig. 42, the optimal NaCl concentration for hybridization as implemented in a dual bead assay is 0.3M NaCl. A detailed description of the experimental procedure used to generate this data is discussed below in Example 4.

5 Turning now to Figs. 43A and 43B, here are bar graph presentations showing the effect of increasing the concentration of $MgCl_2$ in the assay buffer on the dual bead assay sensitivity and an enzyme assay sensitivity, respectively. Data from these figures indicate that a concentration of 30mM $MgCl_2$ in the hybridization buffer is optimal for increasing the signal generated and the assay sensitivity. According
10 to the data shown in Figs. 43A and 43B, the enzyme assay appears to be more sensitive than the dual bead assay in the 30mM $MgCl_2$ treatment. This conclusion is based on the difference in signal within the treatment group from the various target concentrations. Thus as illustrated, the slope of the concentration curve in the 30mM $MgCl_2$ group of the enzyme assay of Fig. 43B is steeper than the
15 corresponding curve in Fig. 43A. Example 5 describes in detail the procedure for carrying out an experiment relating to Fig. 43A.

Referring next to Fig. 44, there is shown a pictorial representation of the use of probe blocking agents to increase the sensitivity of the bead assay. The probe blocking agent used in this particular example is a biotinylated DNA that is
20 complimentary to the probe on the bead. The amount of probe blocking agent used to block excess probes on the bead is such that a pre-determined fraction of probes are blocked by the blocking agent. The use of the probe blocking agent in dual bead assay increases the sensitivity of the assay in that it enhances the probability of target binding to a single capture and reporter bead in a dual bead assay. This
25 may increase the sensitivity of the dual bead assay up to one target per dual bead complex. The use of the biotinylated probe blocking agent allows for the quantitation of the blocking efficiency of the probe blocking agent for optimization of the assay. The amount of biotinylated probes bound to the beads may be quantitated by an enzyme assay using streptavidinated or neutravidinated enzymes
30 including streptavidin-alkaline phosphatase (S-AP) and their appropriate substrates. The choice of enzyme and substrate for use in this test is dictated by the type of detection desired. In general, a colorimetric test is performed wherein the enzyme-substrate reaction produces color that is quantified by a spectrophotometer. Alternatively, streptavidinated or neutravidinated fluorescent tags may also be used

which may be quantified using a fluorimeter or a Fluorimager. Both the colorimetric and fluorescent quantitation may also be carried out using the appropriate optical disc reader as shown in Figs. 1 and 2.

Fig. 45 shows a bar graph presentation of data illustrating the effect of incubation time on the signal generated and the assay sensitivity using different target concentrations during a hybridization reaction in a dual bead assay. The data shows that 2 hours is the minimum incubation time required to generate the maximum signal and sensitivity for the dual bead assay and that a 4 hour or overnight hybridization is not necessary. Example 7, presented below, explains the details regarding the experiment performed to generate the data shown in Fig. 45.

Similarly, Fig. 46 shows a bar graph of data collected illustrating the effect of incubation time and mixing on the hybridization efficiency and the assay sensitivity using different target concentrations in a hybridization reaction as implemented on a dual bead assay. As in Fig. 45, Fig. 46 also shows that 2 hours is an optimal time for hybridization and extending the hybridization time does not increase the signal generated. In addition, mixing significantly increased the hybridization efficiency after 2 hours of hybridization relative to control. Example 7, presented below, explains the details regarding the experiment performed to generate the data shown in Figs. 45 and 46.

Experimental Details

While this invention has been described in detail with reference to the drawing figures, certain examples and further illustrations of the invention are presented below.

EXAMPLE 1

The two-step hybridization method demonstrated in Fig. 12A was used in performing the dual bead assay of this example.

A. Dual Bead Assay

In this example, the dual assay is carried out to detect the gene sequence DYS that is present in male but not in female. The assay is comprised of 3 μ magnetic and capture beads coated with covalently attached capture probe; 2.1 μ fluorescent reporter beads coated with a covalently attached sequence specific for the DYS gene, and target DNA molecule containing DYS sequences. The target

DNA is a synthetic 80 oligonucleotide sequence. The capture probe and reporter probes are 40 nucleotides in length and are complementary to DYS sequence but not to each other.

- The specific methodology employed to prepare the assay involved treating
- 5 1×10^7 capture beads and 2×10^7 reporter beads in 100 microgram per milliliter Salmon Sperm DNA for 1 hr. at room temperature. This pretreatment will reduce non-covalent binding between the capture and reporter beads in the absence of target DNA as shown in Fig. 38. The capture beads were concentrated magnetically with the supernatant being removed. A 100 microliter volume of the
 - 10 hybridization buffer (0.2 M NaCl, 1 mM EDTA, 10 mM $MgCl_2$, 50 mM Tris HCl, pH 7.5, and 5X Denhart's mixture, 10 microgram per milliliter denatured salmon sperm DNA) were added to the capture beads and the beads were re-suspended. Various concentrations of target DNA ranging from 1, 10, 100, 1000 femtomoles were added while mixing at $37^\circ C$ for 2 hours. The beads were magnetically concentrated and
 - 15 the supernatant containing target DNA was removed. A 100 microliter volume of wash buffer (145mM NaCl, 50 mM Tris pH 7.5, 0.1% SDS, 0.05 % Tween, 0.25 % NFDM, 10 mM EDTA) was added and the beads were re-suspended. The beads were magnetically concentrated and the supernatant was again removed. The wash procedure was repeated twice.
 - 20 A 2×10^7 amount of reporter beads in 100 microliter hybridization buffer (0.2 M NaCl, 1 mM EDTA, 10 mM $MgCl_2$, 50 mM Tris HCl, pH 7.5, and 5X Denhart's mixture, 10 microgram per milliliter denatured salmon sperm DNA) were then added to washed capture beads. The beads were re-suspended and incubated while mixing at $37^\circ C$ for an additional 2 hours. After incubation the capture beads were
 - 25 concentrated magnetically, and the supernatant containing unbound reporter beads were removed. A 100 microliter volume of wash buffer (145mM NaCl, 50 mM Tris pH 7.5, 0.1% SDS, 0.05 % Tween, 0.25 % NFDM, 10 mM EDTA) was added and the beads were re-suspended. The beads were magnetically concentrated and the supernatant was again removed. The wash procedure was repeated twice.
 - 30 After the final wash, the beads were re-suspended in 20 microliters of binding buffer (50 mM Tris, 200mM NaCl, 10 mM $MgCl_2$, 0.05% Tween 20, 1% BSA). A 10 microliter volume was loaded on to the disc that was prepared as described below in Part B of this example.

B. Preparation of the Disc

A gold disc was coated with maleic anhydride polystyrene. An amine DNA sequence complementary to the reporter probes (or capture agent) was immobilized on to the discrete reaction zones on the disc. Prior to sample injection, the channels
 5 were blocked with a blocking buffer (50mM Tris, 200mM NaCl, 10 mM MgCl₂, 0.05% Tween 20, 1% BSA, 1% sucrose) to prevent non-covalent binding of the dual bead complex to the disc surface. A perspective view of the disc assembly showing capture agents 220, the capture zones 170, and fluidic circuits as employed in the present invention is illustrated in detail in Figs. 25A-25D. Alternatively, if the
 10 reporter beads are coated with Streptavidin, a capture zone could be created with the capture agent such as BSA Biotin which could be immobilized on to the disc (pretreated with Polystyrene) by passive absorption. A perspective view of the disc assembly showing the use of biotin capture agents is presented in Figs. 26A-26D. Various methods for use in this type of anchoring of beads onto the disc are also
 15 shown in Figs. 15A-15B, 17, 19A-19C, and 23A-23B.

C. Capture of Dual Bead Complex Structure on the Disc

A 10 microliter volume of the dual bead mixture prepared as described in Part A above was loaded in to the disc chamber and the injection ports were sealed. To
 20 facilitate hybridization between the reporter probes on the reporter beads and the capture agents, the disc was centrifuged at low speed (less than 800 rpm) upto 15 minutes. The disc was read in the CD reader at the speed 4X (approx. 1600 rpm) for 5 minutes. Under these conditions, the unbound magnetic capture beads were centrifuged away from the capture zone. The magnetic capture beads that were in
 25 the dual bead complex remained bound to the reporter beads in the capture zone. The steps involved in using the disc to capture and analyze dual bead complexes are presented in detail in Figs. 25A-25D, 26A-26D, and 27A-27D.

D. Quantification of the Dual Bead Complex Structures

30 The amount of target DNA captured could be enumerated by quantifying the number of capture magnetic beads and the number of reporter beads since each type of bead has a distinct signature.

EXAMPLE 2

A. *Dual Bead Assay Multiplexing*

In this example, the dual bead assay is carried out to detect two DNA targets simultaneously. The assay is comprised of 3 μ magnetic capture bead. One population of the magnetic capture bead is coated with capture probes 1 which are complementary to the DNA target 1, another population of magnetic capture beads is coated with capture probes 2 which are complementary to the DNA target 2. Alternatively two different types of magnetic capture beads may be used. There are two distinct types of reporter beads in the assay. The two types may differ by chemical composition (for example Silica and Polystyrene) and/or by size. Various combinations of beads that may be used in a multiplex dual bead assay format are depicted in Fig. 32. One type of reporter bead is coated with reporter probes 1, which are complementary to the DNA target 1. The other reporter beads are coated with reporter probes 2, which are complementary to the DNA target 2. Again the capture probes and the reporter probes are complementary to the respective targets but not to each other.

The specific methodology employed to prepare the dual bead assay multiplexing involved treating 1x10⁷ capture beads and 2x10⁷ reporter beads in 100 μ g/ml salmon sperm DNA for 1 hour at room temperature. This pretreatment will reduce non-covalent binding between the capture and reporter beads in the absence of target DNA. The capture beads were concentrated magnetically with the supernatant being removed. A 100 microliter volume of the hybridization buffer (0.2 M NaCl, 1 mM EDTA, 10 mM MgCl₂, 50 mM Tris HCl, pH 7.5, and 5X Denhart's mixture, 10 microgram per milliliter denatured salmon sperm DNA) were added and the beads were re-suspended. Various concentrations of target DNA ranging from 1, 10, 100, 1000 femto moles were added to the capture beads suspension. The suspension was incubated while mixing at 37°C for 2 hours. The beads were magnetically concentrated and the supernatant containing target DNA was removed. A 100 microliter volume of wash buffer (145mM NaCl, 50 mM Tris pH 7.5, 0.1% SDS, 0.05 % Tween, 0.25 % NFDM, 10 mM EDTA) was added and the beads were re-suspended. The beads were magnetically concentrated and the supernatant was again removed. The wash procedure was repeated twice.

A 2×10^7 amount of reporter beads in 100 microliter hybridization buffer (0.2 M NaCl, 1 mM EDTA, 10 mM $MgCl_2$, 50 mM Tris HCl, pH 7.5, and 5X Denhart's mixture, 10 microgram per milliliter denatured salmon sperm DNA) were then added to washed capture beads. The beads were re-suspended and incubated while
 5 mixing at $37^\circ C$ for an additional 2 hours. After incubation the capture beads were concentrated magnetically, and the supernatant containing unbound reporter beads were removed. A 100 microliter volume of wash buffer (145mM NaCl, 50 mM Tris pH 7.5, 0.1% SDS, 0.05 % Tween, 0.25 % NFDM, 10 mM EDTA) was added and the beads were re-suspended. The beads were magnetically concentrated and the
 10 supernatant was again removed. The wash procedure was repeated twice.

After the final wash, the beads were re-suspended in 20 microliters of binding buffer (50 mM Tris, 200mM NaCl, 10 mM $MgCl_2$, 0.05% Tween 20, 1% BSA). A 10 microliter volume of this solution was loaded on to the disc that was prepared as described in below in Part B of this example.

15

B. Disc Preparation

A gold disc was coated with maleic anhydride polystyrene as described. Distinct reaction zones were created for two types of reporter beads. Each reaction zone consisted of amine DNA sequences complementary to the respective reporter
 20 probes (or capture agents). Prior to sample injection, the channel were blocked with a blocking buffer (50mM Tris, 200mM NaCl, 10 mM $MgCl_2$, 0.05% Tween 20, 1% BSA, 1% sucrose) to prevent non-covalent binding of the dual bead complex to the disc surface. Alternatively, magnetic beads employed in a multiplexing dual bead assay format may be detected using a magneto-optical disc and drive. The
 25 chemical reaction zones, in the magnetic disc format, are replaced by distinctly spaced magnetic capture zones as discussed in conjunction with Fig. 37.

C. Capture of Dual Bead Complex Structure on the Disc

A 10 microlitre volume of the dual bead mixture prepared as described above
 30 in Part A of this example, was loaded in to the disc chamber and the injection ports were sealed. To facilitate hybridization between the reporter probes on the reporter beads and the capture agents, the disc was centrifuged at low speed (less than 800 rpm) for up to 15 minutes. The disc was read in the CD reader at the speed 4X (approx. 1600 rpm) for 5 minutes. Under these conditions, the unbound magnetic

capture beads were centrifuged to the bottom of the channels. The reporter beads bound to the capture zone via hybridization between the reporter probes and their complementary agent.

5 *D. Quantification of the Dual Bead Complex Structures*

The amount of target DNA 1 and 2 captured could be enumerated by quantifying the number of the respective reporter beads in the respective reaction zones.

EXAMPLE 3

10 This study was performed to elucidate the assay sensitivity and range of detection of a genetic dual bead assay.

A. Preparation of Capture and Reporter Beads

The beads used in this experiment were magnetic capture beads (3um
15 Spherotech) and yellow-fluorescent reporter beads (1um Polysciences) each covalently conjugated with DNA transport probes and DNA signal probes, respectively. Approximately 1×10^7 capture beads and 2×10^7 reporter beads were used for this experiment. These beads were washed 3X with PBS and resuspended in 1ml water containing 100 $\mu\text{g/ml}$ digested salmon sperm DNA. The bead solutions
20 in the salmon sperm DNA mixture were then incubated for 1 hour at room temperature. After incubation the beads were washed 3x with wash buffer (145mM NaCl, 50mM Tris, pH 7.5, 0.05% Tween) and 1X with hybridization buffer (0.1M NaCl, 10mM MgCl_2 , 1mM EDTA, 50mM Tris, pH 7.5). The beads were then resuspended in hybridization buffer (containing 100 $\mu\text{g/ml}$ of DNA).

25

B. Dual Bead Assay

A serial dilution of DNA target agents were prepared containing: 100 femtomole, 10 femtomole, 1 femtomole, 0.1 femtomole, 0.01 femtomole, and 0 femtomoles (negative control). Equal amounts of capture beads were then mixed
30 with the various solutions of target and incubated at 37°C for 2 hours to let target hybridized to the 5' capture probe on the beads. After incubation, the bead solutions were washed 3X with the wash buffer (145mM NaCl, 50mM Tris, pH 7.5, 0.05% Tween) and 1X with hybridization buffer (0.1M NaCl, 10mM MgCl_2 , 1mM

EDTA, 50mM Tris, pH 7.5), then resuspended in hybridization buffer (containing 100 µg/ml salmon sperm DNA).

Reporter beads were mixed with the capture bead solution and incubated at 37°C for 2 hours in a rotating mixer. After incubation the bead solutions were washed 6X with 0.5 ml of wash buffer (145mM NaCl, 50mM Tris, pH 7.5, 0.05% Tween, 0.1 % SDS, 0.25% NFDM) and resuspended in 250µl water. The number of reporter beads bound were then quantified by a fluorimeter at Excitation = 500nm, Emission = 530nm, Slit = Ex-2, Em-2, and integration time of 0.1 second.

10

EXAMPLE 4

This study was performed to determine the optimal salt concentration in the hybridization buffer for use in a genetic dual bead type assay.

A. Preparation of Capture and Reporter Beads

15

The beads used in this experiment were magnetic capture beads (3µm Spherotech) and yellow-fluorescent reporter beads (2.1µm from Molecular Probes) each covalently conjugated with DNA transport probes and DNA signal probes, respectively. The beads were washed 1X with hybridization buffer (0.1 M NaCl, 10 mM MgCl₂, 1mM EDTA, 50mM tris, pH 7.5). The beads were pretreated with 0.1% CHAPS and salmon sperm DNA for 1 hour at room temp. The beads were then washed 3X with wash buffer (145mM NaCl, 50 mM Tris, pH 7.5, 0.1% SDS, 0.05% Tween, 0.25% NFDM), 1X with respective hybridization buffer (145mM NaCl, 10mM MgCl₂, 1mMEDTA, 100µg/ml salmon sperm DNA, 50 mM Tris, pH 7.5). After washing, the beads were resuspended in hybridization buffer.

25

B. Dual Bead Assay

The magnetic capture beads, prepared in section A, were divided into 24 10ul aliquots. Six sets of aliquots were diluted in hybridization solution containing various concentrations of NaCl (0mM, 145mM, 300mM, and 400mM). The target DNA mixture was made in the hybridization buffer, of varying salt concentrations, in the following concentrations: 100fmole, 10fmole, 1fmole, 0.1fmole, 0.01fmole, 0 femtomole (negative control). The various target solutions were then mixed with their respective bead solutions according to the salt concentration of the hybridization buffer and incubated at 37°C for 2 hours to let target to hybridize to the

30

5' transport probe on the capture beads. After hybridization, the assay solutions were washed 3X with wash buffer containing the appropriate amount of NaCl required for each treatment group (0mM, 0.145mM, 0.3mM, 0.4mM NaCl), 1X with the appropriate hybridization buffer. The beads were then resuspended in their
 5 respective hybridization buffers containing the appropriate amount of NaCl.

A 100 μ l volume of reporter beads in hybridization buffer containing various NaCl concentrations (0mM, 0.145mM, 0.3mM, 0.4mM NaCl), were added to the appropriate assay solution, so that the same concentration of NaCl was maintained within the different treatment groups. These assay solutions were then incubated at
 10 37°C for 2 hours in a rotating mixer. After incubation, the various solutions were then washed 6X with 0.5 ml of wash buffer containing the appropriate amount of NaCl and once with water also containing NaCl at concentrations equal to that in each respective hybridization buffer (0mM, 0.145mM, 0.3mM, 0.4mM NaCl). The various dual bead solutions were then resuspended in 250 μ l water. The number of
 15 reporter beads bound were then quantified using a fluorimeter at Excitation = 500nm, Emission = 530nm, Slit = Ex-2, Em-2 and an integration time of 0.1 second. The results from this assay are shown in above Fig. 42. Detection of the dual bead complex may be carried out using the optical disc system described as described in conjunction with Fig. 2., a magneto-optical disc system, a fluorescent disc system,
 20 or any similar device. Unique signature traces of a dual bead complex collected from an optical disc reader are shown above in Fig. 29B.

EXAMPLE 5

In this case, magnetic capture beads (3 μ m) from Spherotech and yellow-
 25 fluorescent (2.1 μ m) reporter beads from Molecular Probes were evaluated. This study was performed to determine the optimal MgCl₂ concentration in the hybridization buffer for use in a genetic dual bead type assay.

A. Preparation of Capture and Reporter Beads

30 The magnetic capture beads and yellow-fluorescent reporter beads were each covalently conjugated with DNA transport probes and DNA signal probes, respectively. After conjugation, the beads were washed 1X with hybridization buffer (0.1 M NaCl, 10 mM MgCl₂, 1mM EDTA, 50mM Tris, pH 7.5). The beads were pretreated with 100 μ g/ml salmon sperm DNA for 1 hour at room temp. The

beads were then washed 3X with wash buffer (145mM NaCl, 50 mM Tris, pH 7.5, 0.1% SDS, 0.05% Tween, 0.25% NFDM), 1X with hybridization buffer (145mM NaCl, 10mM MgCl₂, 1mM EDTA, 100µg/ml salmon sperm DNA, 50 mM Tris, pH 7.5). After washing, the beads were resuspended in hybridization buffer.

5

B. Dual Bead Assay

The magnetic capture beads, as prepared in Part A, were divided into 24 10ul aliquots. Six sets of aliquots were diluted in hybridization solution containing various concentrations of MgCl₂ (0mM, 10mM, 20mM, and 30mM). The target DNA mixture
10 was made in hybridization buffer containing varying amounts MgCl₂, in the following concentrations: 100fmole, 10fmole, 1fmole, 0.1fmole, 0.01fmole, 0fmole (negative control). The various target solutions were then mixed with their respective bead solutions according to the salt concentration of the hybridization buffer and incubated at 37°C for 2 hours to let target to hybridize to the 5' transport probe on
15 the capture beads. After hybridization, the assay solutions were washed 3X with wash buffer containing the appropriate amount of MgCl₂ required for each treatment group (0mM, 10mM, 20mM, 30mM MgCl₂), 1X with the appropriate hybridization buffer. The beads were then resuspended in their respective hybridization buffers containing the appropriate amount of MgCl₂.

20 A 100µl volume of reporter beads in hybridization buffer containing various MgCl₂ concentrations (0mM, 10mM, 20mM, 30mM MgCl₂), were added to the appropriate assay solution, so that the same concentration of MgCl₂ was maintained within the different treatment groups. These assay solutions were then incubated at 37°C for 2 hours in rotating mixer. After incubation, the various solutions were then
25 washed 6X with 0.5 ml of wash buffer containing the appropriate amount of NaCl and once with water also containing MgCl₂ at concentrations equal to that in each respective hybridization. The various dual bead solutions were then resuspended in 250µl water. The number of reporter beads bound were then quantified by the fluorimeter at Excitation = 500nm, Emission = 530nm, Slit = Ex-2, Em-2, and an
30 integration of 0.1 second. The results from this assay are shown in above Fig. 43A. The dual bead complex may also be quantified using using an optical disc reader as shown in Figs. 1 and 2.

EXAMPLE 6

The following experiment was performed to determine the effect of using a probe blocking agent to reduce the density of probes on beads on the sensitivity of the dual bead assay.

A. Preparation of Capture and Reporter Beads

The magnetic capture beads (3um Spherotech) and yellow-fluorescent reporter beads (2.1 um Polysciences) were each covalently conjugated with DNA transport probes and DNA signal probes, respectively. After conjugation, the beads were washed 1X with hybridization buffer (0.1 M NaCl, 10 mM MgCl₂, 1mM EDTA, 50mM tris, pH 7.5). The beads were pretreated with 100µg/ml salmon sperm DNA for 1 hour at room temp. The beads were then washed 3X with wash buffer (145mM NaCl, 50 mM Tris, pH 7.5, 0.1% SDS, 0.05% Tween, 0.25% NFDM), 1X with hybridization buffer (100mM NaCl, 10mM MgCl₂, 1mMEDTA, 100µg/ml salmon sperm DNA, 50 mM Tris, pH 7.5). After washing, the beads were resuspended in hybridization buffer.

B. Probe Blocking

A biotinylated transport blocking probe was diluted to the following final concentrations: 500pmole, 50pmole, 35pmole, 30pmoles. A 13µl (2x10⁷) volume of magnetic beads were used for each tube (5 tubes total). A 5µl amount of blocking probe, as prepared above, and 32 µl hybridization buffer was added to each tube. The blocking probes and the transport probes were then hybridized for 2 hours at 37°C. After hybridization, the beads were washed 3X with wash buffer (145nM NaCl, 50nM Tris, pH 7.5, 0.05% Tween) and resuspended in 100µl CDB (2% BSA, 50mM Tris-HCl, pH7.5, 145mM NaCl, 1.0mM MgCl₂, 0.1mM ZnCl₂, 0.05% NaN₃). The reporter beads were prepared in a similar fashion using a biotinylated reporter blocking probe.

C. Probe Density Determination on Beads after Blocking Agent Treatment.

An aliquot of each set of bead solution prepared in Part B was incubated with 100µl of S-AP (1420ng/ml) for one hour at 37°C then washed 3X with wash buffer. After washing, a 100µl volume of S-AP substrate, p-nitrophenyl phosphate, at a
 5 concentration of 3.7mg/ml in 0.1M Tris, 2mM MgCl₂, pH 10 was added to the bead solution. After allowing sufficient time for color development, the solution was analyzed using a spectrophotometer (OD @ 405 nm). The amount of blocking probes on the beads was calculated from the absorbance at 405nm.

10 *D. Dual Bead Assay*

The beads, as prepared in Part B, were washed and resuspended in hybridization buffer containing 100µg/ml salmon sperm DNA and 5X Denhart's solution. A solution target DNA mixture was prepared in the hybridization buffer with the following concentrations: 0fmole-control, 10fmole, 1fmole, 0.1fmole, 0.01fmole,
 15 0.001fmole, 0.0001fmole. The target solutions were then mixed with equal amounts of capture and reporter beads and incubated at 37°C for 2 hours. The capture and reporter beads having been blocked with the same amount of probe blocking agent for each set of assay mixture, i.e., add 10ul 10fmole target to 100ul reporter and capture bead solution each having been blocked with 50pmole blocking probe.
 20 After hybridization, the assay solution was washed 3X with wash buffer and 1X with hybridization buffer and resuspended in hybridization buffer (containing 100 µg/ml salmon sperm DNA and 5X Denhart's Mix). The assay solution was concentrated and resuspended in 250µl water. The number of reporter beads bound were then quantified by the fluorimeter at Excitation = 500nm, Emission = 530nm, Slit = Ex-2,
 25 Em-2, and an integration time of 0.1 seconds.

EXAMPLE 7

The following experiment was performed to determine the optimal
 30 hybridization incubation time of a genetic dual bead assay. The results from this experiment are shown above in Figs. 45 and 46.

A. Preparation of Capture and Reporter Beads

The beads used in this experiment were 25 μ l of capture beads (3 μ m carboxylated magnetic particles at a concentration of 1.5×10^7 beads/ μ l) with 5' transport probes attached by covalent conjugation and 400 μ l of reporter beads (2 μ m YF beads at a concentration of 6.6×10^6 beads/ μ l). These beads were washed 3X with PBS and pretreated with 100 μ g/ml salmon sperm DNA and 0.1% CHAPS for 1 hour at room temperature. The beads were then washed 3X with wash buffer (145mM NaCl, 50mM Tris, pH 7.5, 0.05% Tween) and 1X with hybridization buffer (50mM Tris-HCl pH 7.5, 10mM MgCl₂, 0.1M NaCl, 1mM EDTA). The capture beads were then resuspended in 250 μ l hybridization buffer, and the reporter beads in 400 μ l hybridization buffer.

B. Dual Bead Assay

A set of target DNA solutions are made in hybridization solution with the following concentrations: 0 picomole, 1 picomole, 10 picomole, 100 picomole target.

A test sample was prepared containing 10 μ l capture beads, 15 μ l of reporter beads, 1 μ l salmon sperm DNA and 74 μ l target solution all in hybridization buffer. Aliquots of this test sample were analyzed at various incubation times 30min, 1hr, 2hr, 3hr, 4hr, and overnight. One set was incubated at 37° C without mixing and the other set was mixed on a rotating mixer.

The sample aliquots were washed 6X with 0.5ml wash buffer (145mM NaCl, 50mM Tris, pH 7.5, 0.05% Tween, 0.1 % SDS, 0.25% NFDM) and then resuspended in 202 μ l PBS. The number of reporter beads bound were then quantified by the fluorimeter at Excitation = 450nm, Emission = 480nm, Slit = Ex-1.365, Em-1.05, integration time = 0.1 second.

Concluding Statement

While this invention has been described in detail with reference to certain preferred embodiments and technical examples, it should be appreciated that the present invention is not limited to those precise embodiments or examples. Rather, in view of the present disclosure, which describes the current best mode for practicing the invention, many modifications and variations would present themselves to those of skill in the art without departing from the scope and spirit of this

invention. The scope of the invention is, therefore, indicated by the following claims rather than by the foregoing description. All changes, modifications, and variations coming within the meaning and range of equivalency of the claims are to be considered within their scope.